

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: June 20, 2004, 01:40:43 ; Search time 59.875 Seconds  
(without alignments)  
3923.374 Million cell updates/sec

Title: US-09-000-062-7

Perfect score: 494

Sequence: 1 CTCAGGCGAAGAACAGGTAT.....CGCAGATCCGGGATCG 494

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 682709 seqs, 277475446 residues

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

Issued Patents NA: \*  
1: /cgn2\_6/ptodata/2/ina/5A COMB.seq: \*  
2: /cgn2\_6/ptodata/2/ina/5B COMB.seq: \*  
3: /cgn2\_6/ptodata/2/ina/6A COMB.seq: \*  
4: /cgn2\_6/ptodata/2/ina/6B COMB.seq: \*  
5: /cgn2\_6/ptodata/2/ina/PTUS COMB.seq: \*  
6: /cgn2\_6/ptodata/2/ina/backfiles1.seq: \*

Préd. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	494	100.0	494	US-09-000-062-7	Sequence 7, Appli
2	49.6	10.0	7218	US-08-232-463-14	Sequence 14, Appli
3	38.2	7.7	6669	US-10-204-708-5	Sequence 5, Appli
4	37.2	7.5	832	US-09-621-976-2813	Sequence 2813, Ap
5	37.2	7.5	1230025	US-09-198-452A-1	Sequence 1, Appli
6	37	7.5	1193	US-09-372-422A-23	Sequence 23, Appli
7	36.2	7.3	5562	US-10-204-708-63	Sequence 63, Appli
8	36.2	7.3	6866	US-10-204-708-20	Sequence 20, Appli
9	35.8	7.2	1493	US-08-340-820-24	Sequence 24, Appli
10	35.8	7.2	1493	US-08-593-535-24	Sequence 24, Appli
11	35	7.1	10467	US-10-204-708-1	Sequence 1, Appli
12	34.8	7.0	289	US-09-007-005-17	Sequence 17, Appli
13	34.8	7.0	289	US-09-244-796-17	Sequence 17, Appli
14	34.4	7.0	10467	US-10-204-708-2	Sequence 2, Appli
15	34.2	6.9	1406	US-08-913-842-6	Sequence 6, Appli
16	34	6.9	1664976	US-08-916-421B-1	Sequence 1, Appli
17	33.8	6.8	1051	US-09-396-149-15	Sequence 15, Appli
18	33.6	6.8	531	US-09-328-352-2861	Sequence 2861, Ap
19	33.6	6.8	7304	US-10-204-708-43	Sequence 43, Appli
20	33.6	6.8	11015	US-10-204-708-55	Sequence 55, Appli
21	33.2	6.7	1896	US-09-107-532A-894	Sequence 894, App
22	32.8	6.6	5666	US-10-204-708-29	Sequence 29, Appli
23	32.8	6.6	8961	US-10-204-708-80	Sequence 80, Appli
24	32.8	6.6	15071	US-09-358-082A-29	Sequence 29, Appli
25	32.6	6.6	11014	US-08-956-171E-91	Sequence 91, Appli
26	32.4	6.6	1545	US-09-107-532A-2628	Sequence 2628, Ap
27	32.4	6.6	5360	US-10-204-708-65	Sequence 65, Appli

## ALIGNMENTS

### RESULT 1

US-09-000-062-7

; Sequence 7, Application US/09000062

; Patent No. 6338961

; GENERAL INFORMATION:

; APPLICANT: DEROSE, Richard

; APPLICANT: CHAUBET, Nicole

; TITLE OF INVENTION: ISOLATED DNA SEQUENCE CAPABLE OF SERVING AS REGULATORY

; TITLE OF INVENTION: ELEMENT IN A CHIMERIC GENE WHICH CAN BE USED FOR THE

; TITLE OF INVENTION: TRANSFORMATION OF PLANTS

; FILE REFERENCE: 022650-453

; CURRENT APPLICATION NUMBER: US/09/000,062

; CURRENT FILING DATE: 1998-05-29

; EARLIER APPLICATION NUMBER: PCT/FR96/01109

; EARLIER FILING DATE: 1996-07-17

; EARLIER APPLICATION NUMBER: FR 95/08980

; EARLIER FILING DATE: 1995-07-19

; NUMBER OF SEQ ID NOS: 22

; SOFTWARE: PatentIn Ver. 2.0

; SEQ ID NO 7

; LENGTH: 494

; TYPE: DNA

; ORGANISM: Zea mays

; US-09-000-062-7

Query Match 100.0%; Score 494; DB 4; Length 494;

Best Local Similarity 100.0%; Pred. No. 28-128;

Matches 494; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCAGGCGAAGAACAGGTATGTTTGTGTAATTAGATCAGGGTTTAGGTCTTCCAT 60

Db 1 CTCAGGCGAAGAACAGGTATGTTTGTGTAATTAGATCAGGGTTTAGGTCTTCCAT 60

QY 61 TACTTTTATGTTTTTCTGTTACTGTCCTCCGATCTGATTTTACGACAAATAGATTT 120

Db 61 TACTTTTATGTTTTTCTGTTACTGTCCTCCGATCTGATTTTACGACAAATAGATTT 120

QY 121 CGGGTTTGTCCCATCCAGTTTGAAATAAAGTCGCTCTTTTAAAGTTTGCTGGATCGA 180

Db 121 CGGGTTTGTCCCATCCAGTTTGAAATAAAGTCGCTCTTTTAAAGTTTGCTGGATCGA 180

QY 181 TAAACCTGTGAAGTTGAGTCTAGTCGATTTATTTGGATGATCATTCTTCTATCGTTTTT 240

Db 181 TAAACCTGTGAAGTTGAGTCTAGTCGATTTATTTGGATGATCATTCTTCTATCGTTTTT 240

QY 241 TCTTGCTTCAAGTTTCTGTATACAGATTGTCTGTGTGCGATTGTTCATTACTACCG 300

Db 241 TCTTGCTTCAAGTTTCTGTATACAGATTGTCTGTGTGCGATTGTTCATTACTACCG 300

Sequence 41, Appli  
Sequence 71, Appli  
Sequence 244, App  
Sequence 3, Appli  
Sequence 10903, A  
Sequence 21, Appli  
Sequence 10, Appli  
Sequence 14, Appli  
Sequence 400, App  
Sequence 760, App  
Sequence 852, App  
Sequence 7, Appli  
Sequence 171, App  
Sequence 27, Appli  
Sequence 46, Appli  
Sequence 3, Appli  
Sequence 3, Appli  
Sequence 3, Appli

4 US-10-204-708-41  
4 US-10-204-708-71  
4 US-08-356-171E-244  
4 US-09-816-093-3  
4 US-09-621-976-10903  
4 US-09-453-702B-21  
4 US-10-204-708-10  
4 US-10-204-708-14  
4 US-09-134-001C-400  
4 US-09-976-594-760  
4 US-09-134-001C-852  
4 US-09-272-496-7  
4 US-08-956-171E-171  
4 US-10-204-708-27  
4 US-09-755-665-46  
4 US-08-307-382-3  
4 US-08-366-779-3  
4 US-08-478-727-3

6.5 8537  
6.5 8607  
6.5 2022  
6.5 46718  
6.4 476  
6.4 4998  
6.4 6070  
6.4 6113  
6.4 789  
6.4 1453  
6.4 2301  
6.4 4072  
6.4 11126  
6.4 11131  
6.4 41100  
6.4 1884  
6.4 1884  
6.4 1884

28 32.2  
29 32.2  
30 32  
31 32  
32 31.8  
33 31.8  
34 31.8  
35 31.8  
36 31.6  
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38 31.6  
39 31.6  
40 31.6  
41 31.6  
42 31.6  
43 31.4  
44 31.4  
45 31.4

301 TGATCGAGAACTAGGTTTCAGAGTCAATTTGCCCCCTTTTGGTTATATCTGGTTCGAT 360  
Db 301 TGATCGAGAACTAGGTTTCAGAGTCAATTTGCCCCCTTTTGGTTATATCTGGTTCGAT 360  
361 AACGATTCACCTGATAGGTTTAAAGTGTGACGTTTAGTATTCCTTCAAAA 420  
Db 361 AACGATTCACCTGATAGGTTTAAAGTGTGACGTTTAGTATTCCTTCAAAA 420  
421 TTTAGTTATGATATGAATGAAATCCCGAATGACGTGTTCAATTTCTTTAAATGCGCAGA 480  
Db 421 TTTAGTTATGATATGAATGAAATCCCGAATGACGTGTTCAATTTCTTTAAATGCGCAGA 480  
481 TCCCGGATCTGCG 494  
Db 481 TCCCGGATCTGCG 494

RESULT 2  
US-08-232-463-14  
; Sequence 14, Application US/08232463  
; Patent No. 5670367  
; GENERAL INFORMATION:  
; APPLICANT: DORNER, F.  
; APPLICANT: SCHEIFLINGER, F.  
; APPLICANT: FALKNER, F. G.  
; TITLE OF INVENTION: RECOMBINANT FOMLPOX VIRUS  
; NUMBER OF SEQUENCES: 52  
; CORRESPONDENCE ADDRESS:  
; ADDRESS: Foley & Hardner  
; STREET: 1800 Diagonal Road, Suite 500  
; CITY: Alexandria  
; STATE: VA  
; COUNTRY: USA  
; ZIP: 22313-0299  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/232,463  
; FILING DATE:  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US/07/935,313  
; FILING DATE:  
; APPLICATION NUMBER: EP 91 114 300.6  
; FILING DATE: 26-AUG-1991  
; ATTORNEY/AGENT INFORMATION:  
; NAME: BENT, Stephen A.  
; REGISTRATION NUMBER: 29,768  
; REFERENCE/DOCKET NUMBER: 30472/114 IMMU  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (703)836-9300  
; TELEFAX: (703)683-4109  
; TELEX: 899149  
; INFORMATION FOR SEQ ID NO: 14:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 7218 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; IMMEDIATE SOURCE:  
; CLONE: pTZ9pt-Fls  
US-08-232-463-14

Query Match 10.0%; Score 49.6; DB 1; Length 7218;  
Best Local Similarity 6.1%; Pred. No. 0.00022;  
Matches 25; Conservative 213; Mismatches 172; Indels 0; Gaps 0;  
QY 14 CAGGTATGATTTCTTTGTAATTAGATCAGGGGTTTAGGCTTTCCATTACTTTTAAATGT 73  
Db 1046 CAGGTACGAGGAGCTTCCGATTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT 1105

74 TTTTCTCTTACTGTCCTCCGCACTGATTTTACGACAATAGAGTTTCGGGTTTGTCCC 133  
Db 1106 YY 1165  
134 ATTCAGTTTGAATAAATCGTCCGCTCTTTTAAAGTTTCTCGATCGATAAACCCTGTGAAG 193  
Db 1166 YY 1225  
194 ATTGAGTCTAGTCGATTTATTTGGATGATCCATCTTCTCATCGTTTCTTCTGCTTCGAAG 253  
Db 1226 YY 1285  
254 TTCTGTATAACCAAGATTTGTCTGTGTGGGATGTGTCATTTACCTAGCCGTGTATCGAAGACT 313  
Db 1286 YY 1345  
314 AGGTTTTCGAGTCAATTTTGCCCTTTTGGTATATCTGTTTCGATACGATTCATCTG 373  
Db 1346 YY 1405  
374 GATTAGGTTTAAAGTGTGACGTTTAGTATTTCCAAATTTCTTCAAAATTT 423  
Db 1406 YY 1455

RESULT 3  
US-10-204-708-5  
; Sequence 5, Application US/10204708  
; Patent No. 5677731  
; GENERAL INFORMATION:  
; APPLICANT: OLEK, Alexander  
; APPLICANT: PIEPENBROCK, Christian  
; APPLICANT: BERLIN, Kurt  
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication  
; FILE REFERENCE: 5013.1012  
; CURRENT APPLICATION NUMBER: US/10/204,708  
; PRIOR FILING DATE: 2003-05-06  
; PRIOR APPLICATION NUMBER: PCT/EP01/03971  
; PRIOR FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: DE 10019058.8  
; PRIOR FILING DATE: 2000-04-06  
; PRIOR APPLICATION NUMBER: DE 10019173.8  
; PRIOR FILING DATE: 2000-04-07  
; PRIOR APPLICATION NUMBER: DE 10032529.7  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: DE 10043826.1  
; PRIOR FILING DATE: 2000-09-01  
; NUMBER OF SEQ ID NOS: 98  
; SEQ ID NO 5  
; LENGTH: 6669  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)  
US-10-204-708-5

Query Match 7.7%; Score 38.2; DB 4; Length 6669;  
Best Local Similarity 48.0%; Pred. No. 0.32;  
Matches 109; Conservative 0; Mismatches 118; Indels 0; Gaps 0;  
QY 50 GGTCTTTCCATTACTTTTAAATGTTTTTCTGTTACTGTCTCCGCACTCTGATTTCGA 109  
Db 5179 GGTCTTTGATTTTTTTTTTTTTTTTATTTATTTTGTGTTTTTTCGGGTGTTTAC 5238  
QY 110 CAATAGAGTTTCGGGTTTCTGCCATTCAGTTTGAATAAAGTCGCTTTTAAGTT 169  
Db 5239 GGATTTGGTTAAATTAATTTTTTTTTTTTAAAGTTTGAATGATTTGTTTTTGGTA 5298  
QY 170 TGCTGGATCGATAAACCTGTGAGAGATTGAGTCTAGTCCATTTTATGATGATCCATCTT 229  
Db 5299 TTTTCTTTGCTGTTATTTTAGGTTTAGATTTAATACGGATTTGATTAGTATTTTT 5358

QY 230 CATCGTTTTTCTTGCTTCTGAAGTCTCTGATAAACGAGATTTGTCG 276  
|||||  
nb 5359 TGAATTTTTTTTCGATTATAGTATTTTACGTAGCGAGTATTATTG 5405

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RESULT 4
US-09-621-976-2813
; Sequence 2813, Application US/09621976
; Patent No. 8639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jober, S.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: ESTs and Encoded Human
; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621.976
; CURRENT FILING DATE: 2000-07-21
; NUMBER OF SEQ ID NOS: 19335
; SOFTWARE: Patent.pm
; SEQ ID NO 2813
; LENGTH: 832
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 235..399
US-09-621-976-2813

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Query Match	7.5%;	Score 37.2;	DB 4;	Length 832;
Best Local Similarity	9.6%;	Pred. No. 0.31;		
Marches	35.	Conservative	176;	Mismatches 154;
				Indels 1;
				Gaps 1;

Qy	71	TGTTTTTCTGTTACTGCTCTCGCGGATCTGATTTTACGCAATAGAGTTTCTCGGGTTTTGT	130
Db	1	YRWYKTTTYYAKGKTKWKSWSYTWYKYYKTYWRWRKKKKAUWTKWTWTWYW	60
Qy	131	CCCATTCCAGTTTGAANAATAACAGTCGCTCTTTTAAAGTTTGTGGATCGATAAACCTGTG	190
Db	61	RYAMWGTYYKKAMCTKTKKKKKGYMMWWTWGRHSYNAMWTRTWGAYYRSMYWWR	120
Qy	191	AAGATTGAGTCTAGTCGGAATTATTGATGATGCATCTCTTCATCGTTTTTTTCTGCTTCG	250
Db	121	YRCWKXKAYVRKTTTCYSKSGMTWWRKKKAWTTWWKKTYYAAATRYWMMCMWTKRFAAS	180
Qy	251	AAGTTCGTGATAACACAGATTGTCTGTGTGGAGTTGTCATTACCTACGTAGCGGTGATCGAGA	310
Db	181	WWYCWGWKARKWSTWRSRSYASARSACRCCYCSGWSGMSKYYWWRWRGWATGAGM	240
Qy	311	ACTAGGGTTTTCGAGTCAATTTTGGCCCTTTTGGTTATATCTGGTTCGATACGATTTCAT	370
Db	241	KAWRASCMWRKYAGSKTYSKSMCMWTRSWKYCYTKABWTGYCYRKGGMWKGGRWY	300
Qy	371	-CTGAGATTAGGGTTTAACTGTGTGACGTTTAGTATTCCAAATTTCTTCAAAATTTTAGTTAT	429
Db	301	ASKKYMWKEWWNCABMYRYSYTGTRASMMWRWRYTYTMMKWKYAWASAAERWAAWNAW	360
Qy	430	GGATAA	435
Db	361	RRACAA	366

RESULT 5  
US-09-198-452A-1  
; Sequence 1, Application US/09198452A  
; Patent No. 6553294  
; GENERAL INFORMATION:  
; APPLICANT: Griffrats, R.  
; TITLE OF INVENTION: Chlamydia pneumoniae genomic sequence and polypeptides, fragments  
; TITLE OF INVENTION: thereof and uses thereof, in particular for the diagnosis, prevention  
; TITLE OF INVENTION: and treatment of infection  
; FILE REFERENCE: 9710-003-999  
; CURRENT APPLICATION NUMBER: US/09/198.452A

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CURRENT BILLING DATE: 1998-11-24
NUMBER OF SEQ ID NOS: 6849
SEQ ID NO 1
LENGTH: 1230025
TYPE: DNA
ORGANISM: Chlamydia pneumoniae
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)..(15000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (15001)..(30000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (30001)..(45000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (45001)..(60000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (60001)..(75000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (75001)..(90000)
OTHER INFORMATION: n=a or c or g or t
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NAME/KEY: misc_feature
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OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (300001)..(315000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc_feature
LOCATION: (315001)..(330000)
OTHER INFORMATION: n=a or c or g or t

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LOCATION: (690001)..(705000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc feature
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OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc feature
LOCATION: (720001)..(735000)
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LOCATION: (885001)..(900000)
OTHER INFORMATION: n=a or c or g or t
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LOCATION: (900001)..(915000)
OTHER INFORMATION: n=a or c or g or t
NAME/KEY: misc feature

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QY	34	TTAGATCAGGGGTTTAGGTCCTTCCATTACTCTTTAAATGTTTTCTGTACTGTCTCG	93
Db	1129940	TGAGTAGAGAGAAATGTTNATTTCTTCTNNCAANAATTTATCTTACCAGTCTCT	11299599
QY	94	CGATCTGATTTTACGACAATAGAGTTTCGGGTTTGTCCCATTCAGTTTGAAAAATAAC	153
Db	1130000	CTATTTGTCTGAGTTACTTGAGTGATGCGTGAATTTTACATGTGACTTTTGGATCATGG	1130059
QY	154	GTCGGCTCTTTTAAGTTTGTGCGATCGATATAAAGCTGTGAAGATTGACTCTAGTCGATTTAT	213
Db	1130060	ATAACTGGAACTTTTATAAAGATATCGAGTTTCAGAAAAGAGAAAAAACCTCTTTTAT	1130119
QY	214	TGATGATCCATCTTCATCGTTTTTTTTCTGCTCGAAGTTCTGTATATACCAGATTTGT	273
Db	1130120	TTAATGATTTATTTTTTAAAGTATATAAATAATTTATATATAAATAATTTTACITTTTT	1130179
QY	274	CTGTGTGCGATTTGCTATTACCTAGCCGTGTATCGAGAACTAGGGTTTTCGAGTCAATTTT	333
Db	1130180	AATAATTAATAAATGATAAATGTTTGCACCTTATAATAAATATAAATTATAGATTTTGATGC	1130239
QY	334	GCCCCCTTTTGGTTATATTC	351
Db	1130240	TTGTCTATTCTTTTGTGATC	1130257

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RESULT 8
US-10-204-708-20
; Sequence 20, Application US/10204708
; Patent No. 6677731
; GENERAL INFORMATION:
; APPLICANT: OLEK, Alexander
; APPLICANT: PIEPENBROCK, Christian
; APPLICANT: BERLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
; TITLE OF INVENTION: by Assessing DNA Methylation
; FILE REFERENCE: 5013.1012
; CURRENT APPLICATION NUMBER: US/10/204,708
; CURRENT FILING DATE: 2003-05-06
; PRIOR APPLICATION NUMBER: PCT/EP01/03971
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: DE 10019058.8
; PRIOR FILING DATE: 2000-04-06
; PRIOR APPLICATION NUMBER: DE 10019173.8
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: DE 10032529.7
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: DE 10043826.1
; PRIOR FILING DATE: 2000-09-01
; NUMBER OF SEQ ID NOS: 98
; SEQ ID NO 20
; LENGTH: 6866
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-204-708-20

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Query Match 7.3%; Score 36.2; DB 4; Length 6866;  
Best Local Similarity 45.3%; Pred. No. 1.2;  
Matches 208; Conservative 0; Mismatches 248; Indels 3; Gaps 2;  
10 AGAACAGGTATGATTTTGTGTAATTAGATCATCGGGTTAGGCTTTCCTTACTTTTAA 69

Query Match	7.5%	Score 37;	DB 4;	Length 1193;
Best Local Similarity	53.9%;	Pred. No. 0.4;		
Matches 76;	Conservative	0;	Mismatches 65;	Indels 0;
Gaps	0;			

  

18	TAAGATTGTTGTTGAATTAGATCAGGGGTTTAGGTCTTTCCATTACTTTTAAATGTTTT	77
QY		
1182	TT	1123
DBb		
78	TCTGTTACTCTCCGCGATCTGATTTTACGACAATAGAGTTTCGGGTTTTGTCGCCATTC	137
QY		
1122	TTCGAAACGAAATCAGCTGTTTTTCCCACTC	1063
DBb		
138	CAGTTTGAATAATAACGTCGC	158
QY		
1062	CAAGCGCATCGAAACGCGCAG	1042
DBb		

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RESULT 7
US-10-204-708-63
  / Sequence 63, Application US/10204708
  / Patent No. 6677731
  / GENERAL INFORMATION:
  / APPLICANT: OLEK, Alexander
  / APPLICANT: PIEPNBROCK, Christian
  / APPLICANT: BEBLIN, Kurt
  / TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
  / TITLE OF INVENTION: by Assessing DNA Methylation
  / FILE REFERENCE: 5013.1012
  / CURRENT APPLICATION NUMBER: US/10/204,708
  / CURRENT FILING DATE: 2003-05-06
  / PRIOR APPLICATION NUMBER: PCT/EP01/03971
  / PRIOR FILING DATE: 2001-04-06
  / PRIOR APPLICATION NUMBER: DE 10019058.8
  / PRIOR FILING DATE: 2000-04-06
  / PRIOR APPLICATION NUMBER: DE 10019173.8
  / PRIOR FILING DATE: 2000-04-07
  / PRIOR APPLICATION NUMBER: DE 10032529.7
  / PRIOR FILING DATE: 2000-06-30
  / PRIOR APPLICATION NUMBER: DE 10043826.1
  / PRIOR FILING DATE: 2000-09-01
  / NUMBER OF SEQ ID NOS: 98
  / SEQ ID NO 63
  / LENGTH: 5562
  / TYPE: DNA
  / ORGANISM: Artificial Sequence
  / FEATURE:
  / OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)

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PRIOR APPLICATION DATA:  
APPLICATION NUMBER: JP 07-073043  
FILING DATE: 30-MAR-1995  
PRIOR APPLICATION DATA: PCT/JP96/00777  
APPLICATION NUMBER: PCT/JP96/00777  
FILING DATE: 26-MAR-1996  
ATTORNEY/AGENT INFORMATION:  
NAME: BROWDY, Roger L.  
REGISTRATION NUMBER: 25,618  
REFERENCE/DOCKET NUMBER: OHBA-1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (202) 628-5197  
TELEFAX: (202) 737-3528  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1406 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: CDNA  
US-08-913-842-6

Query Match 6.9%; Score 34.2; DB 3; Length 1406;  
Best Local Similarity 53.3%; Pred. No. 2.5;  
Matches 72; Conservative 0; Mismatches 63; Indels 0; Gaps 0;

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Qy	399	TAGTATTCCAATTTCTTCAAAATTTAGTTATGGATAATGAAATCCCGAATTGACTGTC	458
Db	1058	TTTGGTGTATTTTATATAATTTTAAATTTCTTTATATAAATATGACTCTTCGTTG	999
Qy	459	AATTCTTTGTTAAAT	473
Db	998	ATTCAAAATTTAAAT	984

Search completed: June 20, 2004, 05:03:44  
Job time : 73.875 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: June 20, 2004, 00:55:23 ; Search time 2930.58 Seconds  
(without alignments)  
6182.178 Million cell updates/sec

Title: US-09-000-062-6  
Perfect score: 418  
Sequence: 1 tgaagtagattcttcgatcc.....taattgtgaacagatccc 418

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 3470272 seqs, 21671516995 residues

Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl:

- 1: gb\_ba.\*
- 2: gb\_htg.\*
- 3: gb\_in.\*
- 4: gb\_ov.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sy.\*
- 13: gb\_un.\*
- 14: gb\_vt.\*
- 15: em\_ba.\*
- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
- 19: em\_mu.\*
- 20: em\_ov.\*
- 21: em\_or.\*
- 22: em\_ov.\*
- 23: em\_pat.\*
- 24: em\_ph.\*
- 25: em\_pl.\*
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- 27: em\_sts.\*
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- 30: em\_htg\_hum.\*
- 31: em\_htg\_inv.\*
- 32: em\_htg\_other.\*
- 33: em\_htg\_mus.\*
- 34: em\_htg\_pln.\*
- 35: em\_htg\_rtd.\*
- 36: em\_htg\_mam.\*
- 37: em\_htg\_vrt.\*
- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	418	100.0	418	6	A59349 Sequence 6
2	418	100.0	418	6	AR182671 Sequence
3	415	99.3	4833	8	ATH3G
4	411.8	98.5	112067	8	ATCHRIY92
5	411.8	98.5	112067	8	ATTSJ17
6	54.4	13.0	5998	6	AX345536 Sequence
7	52	12.4	349980	6	AX344561 Sequence
8	51.2	12.2	253151	3	AE014842 Plasmid
9	50.6	12.1	17527	6	AX339160 Sequence
10	50.6	12.1	17527	6	AX346334 Sequence
11	50.2	12.0	6306	6	AX278060 Sequence
12	50.2	12.0	6306	6	AX323843 Sequence
13	49.6	11.9	206653	2	EX470257
14	49.2	11.8	127172	2	AC141780
15	49.2	11.8	349980	6	AX344573 Sequence
16	49.2	11.8	349980	6	AX344574 Sequence
17	49	11.7	5987	6	AX346464 Sequence
18	49	11.7	5987	6	AX458585 Sequence
19	49	11.7	5987	6	AX795831 Sequence
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21	49	11.7	5987	6	AX822343 Sequence
22	49	11.7	5987	6	AX822471 Sequence
23	49	11.7	5987	6	AX825983 Sequence
24	49	11.7	5987	6	AX826111 Sequence
25	48.8	11.7	15479	6	AX348350 Sequence
26	48.4	11.6	17848	6	AX310889 Dictyost
27	48.4	11.6	17848	6	AX277865 Sequence
28	48.4	11.6	17848	6	AX323550 Sequence
29	48.4	11.6	17848	6	AX348363 Sequence
30	48	11.5	2501	6	AX598874 Sequence
31	48	11.5	2501	6	AX599020 Sequence
32	48	11.5	6059	6	AX346382 Sequence
33	48	11.5	6245	6	AX251202 Sequence
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42	47.6	11.4	5649	6	AX822418 Sequence
43	47.6	11.4	5649	6	AX826058 Sequence
44	47.4	11.3	21537	6	AX346901 Sequence
45	47.4	11.3	349980	6	AX344563 Sequence

ALIGNMENTS

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LOCUS A59349 418 bp DNA linear PAT 06-MAR-1998  
DEFINITION Sequence 6 from Patent WO9704114.  
ACCESSION A59349  
VERSION A59349.1 GI:3714675  
KEYWORDS  
SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eurosids II; Brassicaceae; Arabidopsis.  
REFERENCE 1  
AUTHORS Derose,R., Chaubet,N. and Gigot,C.  
TITLE ISOLATED DNA SEQUENCE FOR USE AS A REGULATOR REGION IN A CHIMERIC

```
GENE USEFUL FOR TRANSFORMING PLANTS
Patent: WO 9704114-A 6 06-FEB-1997;
RHONE POULENC AGROCHIMIE (FR)
Other publication FR 2736929 970124.
Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 2.le-81;
Matches 418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAGGTACGATCTTCGATCCCTCTTGGATTTCTCGGAATATATTTTCGGTGATCGTGA 60
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QY 61 AACTACTCGATCGCTCGATAGGTGGTACGAAATAGGCGAGATTAGTTCTATTCTTGG 120
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DEFINITION A.thaliana H3 gene 1 and H3 gene 2 for H3.3-like histone variant.
ACCESSION X60429
VERSION X60429.1 GI:16323
KEYWORDS histone; histone H3; histone H3.3 homologue.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE 1 (bases 1 to 4833)
AUTHORS Chaubet,N. Clement,B. and Gigot,C.
TITLE Genes encoding a histone H3.3-like variant in Arabidopsis contain
intervening sequences
JOURNAL J. Mol. Biol. 225 (2), 569-574 (1992)
MEDLINE 92277663
PUBMED 1593639
REFERENCE 2 (bases 1 to 4833)
AUTHORS Gigot,C.
TITLE Direct Submission
JOURNAL Submitted (17-SEP-1991) C. Gigot, Inst de Biologie Mol des Plantes,
12 Rue du General Zimmer, 67084 Strasbourg Cedex, FRANCE
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Query Match 100.0%; Score 418; DB 6; Length 418;
Best Local Similarity 100.0%; Pred. No. 2.le-81;
Matches 418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 121 CCATTATCTTGTTCCTCGCGAATGATCTCCGATATAGGATTTAGTTTCTATTCTTGG 180
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DB 301 TTTCTTGTATTCGCGATTCGCAATAGGATTTCTTTGGTTTGTGTTGATCTTACGATA 360
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RESULT 2
LOCUS AR182671 418 bp DNA linear PAT 20-APR-2002
DEFINITION Sequence 6 from patent US 6338961.
ACCESSION AR182671
VERSION AR182671.1 GI:20225878
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 418)
AUTHORS DeRose,R. Chaubet,N. and Gigot,C.
TITLE Isolated DNA sequence capable of serving as regulatory element in a
chimeric gene which can be used for the transformation of plants
JOURNAL Patent: US 6338961-A 6 15-JAN-2002;
MEDLINE
PUBMED
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Best Local Similarity 100.0%; Pred. No. 2.le-81;
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Best Local Similarity 100.0%; Pred. No. 7.1e-81;  
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QY 688 TGAGGTACGATCTTCGATCCCTCTTTGATTTTCCTCGAAATATTTTCGGTGATCGCA 747  
Db |||||  
QY 61 AACTACTGGAATCGCTCGATAGGTGGTACGAATAGGCGAGATTAGTTCTTATCTTGG 120  
Db |||||  
QY 748 AACTACTGGAATCGCTCGATAGGTGGTACGAATAGGCGAGATTAGTTCTTATCTTGG 807  
Db |||||  
QY 121 CCATTATCTGTTTCTTCGCGGAATGATCTTCGATATAAGATTTTAGGTTAGAGATGAA 180  
Db |||||  
QY 808 CCATTATCTGTTTCTTCGCGGAATGATCTTCGATATAAGATTTTAGGTTAGAGATGAA 867  
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RESULT 4  
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LOCUS Arabidopsis thaliana DNA chromosome 4, contig fragment No. 92.  
DEFINITION Arabidopsis thaliana DNA chromosome 4, contig fragment No. 92.  
ACCESSION AL161596  
VERSION AL161596.2 GI:7271037  
KEYWORDS  
SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.  
1 (bases 1 to 107700)  
Murphy,G., Ridley,P., Hudson,S., Mewes,H.W., Lemcke,K. and  
Mayer,K.F.X.  
Unpublished  
2 (bases 9546 to 9977)  
Voickaert,G., Grymonprez,B., Voet,M., Robben,J., Mewes,H.W.,  
Lemcke,K. and Mayer,K.F.X.  
Unpublished  
3 (bases 107578 to 112067)  
Rose,M., Hempel,S., Entian,K.-D., Mewes,H.W., Lemcke,K. and  
Mayer,K.F.X.  
Unpublished  
4 (bases 1 to 112067)  
EU Arabidopsis sequencing project.  
Direct Submission  
Submitted (10-MAR-2000) MIPS, at the Max-Planck-Institut fuer  
Biochemie, Am Klopferspitze 18a, D-82152 Martinsried, FRG, E-mail:

lemcke@mips.biochem.mpg.de, mayer@mips.biochem.mpg.de Project Coordinator: Mike Bevan, Molecular Genetics Department, Cambridge Laboratory, John Innes Centre, Colney Lane, NR4 7UJ Norwich, UK, E-mail: michael.bevan@bbsrc.ac.uk

Information on performance of analysis and a more detailed annotation of this entry and other sequences of chromosomes 3, 4 and 5 can be viewed at: <http://www.mips.biochem.mpg.de/proj/thal/> this fragment has an overlap with ATCHRIV91 at the 5' end and an overlap with ATCHRIV93 at the 3' end.

## COMMENT

## FEATURES

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AA131-151; Multicopper oxidases signatures  
AA540-540; Multicopper oxidases signatures  
AA543-556; Prokaryotic membrane lipoprotein lipid attachment site AA567-577  
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RESULT 5  
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 LOCUS  
 DEFINITION Arabidopsis thaliana DNA chromosome 4, BAC clone T5J17 (BSSA project).  
 ACCESSION AL035708  
 VERSION AL035708.2 GI:5918309  
 KEYWORDS Arabidopsis thaliana (thale cress)  
 SOURCE Arabidopsis thaliana  
 ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

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FSSLSLDLIDLTAVTGILYAWELCSYASPKIFWILAYLGAAGTAIRNFSFSGK  
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Best Local Similarity 99.5%; Pred. No. 2.4e-80;  
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QY 61 AACTACTGGAATCGCTCGATAGGTGTTACGAAATTCGCGAGATTAGTTCTATTCTTGG 120  
DB 96175 AACTACTGGAATCGCTCGATAGGTGTTACGAAATTCGCGAGATTAGTTCTATTCTTGG 96116

QY 121 CCATTATCTTCTTCTTCGCGGAATGATCTTCCGTATATAAGATTTCGTTAGAGATGAA 180  
DB 96115 CCATTATCTTCTTCTTCGCGGAATGATCTTCCGTATATAAGATTTCGTTAGAGATGAA 96056

QY 181 TCGTATACGATAGTTTCATCACCAGATAGTTTCTTCTCTAGAGATCTCTGAAATCTCGA 240  
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QY 241 TAGTTTTACATGCTGTAATAGATTGTTCTTATTCGCGCATGTTGTGATTAGGTTTTGAT 300  
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DB 95935 TTCTTGTATTAGCGATTGCAATTAGGATTTCTTGGTTTCTTGGTTTCTTGGTTTCTTACGATA 95876

QY 361 CATTCCTGCAATTAAGTATGATGATCAATTAATCTTGTAAATTTCTTGAACAGAT 415  
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 Db 95875 CATTCGGCAATTAAGTATGATGATCAATTAATCTTGTAAATTTCTTGAACAGAT 95821  
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RESULT 6  
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 DEFINITION Sequence 1607 from Patent WO0200928.  
 ACCESSION AX344561  
 VERSION AX344561.1 GI:18492442  
 KEYWORDS synthetic construct  
 SOURCE synthetic construct  
 ORGANISM artificial sequences.

REFERENCE  
 1  
 Olek.A., Piepenbrock,C. and Berlin,K.  
 TITLE Diagnosis of diseases associated with the immune system  
 JOURNAL Patent: WO 0200928-A 1607 03-JAN-2002;  
 EpiGenomics AG (DE)  
 FEATURES  
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 Best Local Similarity 49.7%; Pred. No. 0.045;  
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 78 GATAGTGTGACAAATTAAGCGAGATTAATCTTCTTATCTTGGCCATTAATCTTGTCTT 137  
 4382 GAGTAGTGGTTTGAGTATAGTAAATGTAGTTTCGAATTTTGGGTTTAAATGATTTT 4441  
 138 CGCCGATGATCTCCGTA- TAAAGATTTTAGTTAGAGATGAATCGTAGCTAGATT 196  
 4442 TGTTTAGTTTGTAGTTAGTTAGGATTAAGTTATAGTTATGTTTGTAGTTAATAA 4501  
 197 CATCACCAGATAGTTCTTTGTCTAGAAATCTCTGAAATCTCGATAGTTTTCACATGTT 256  
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 4562 TTTTGAATTTTGGTTTAGCGGATTTTATTTTATTTTGTAGTTTGTGTTATATA 4621  
 317 TTGCAATTAGGGATTTTCTTTGTTTGTGTTGATCTTACGATACATTCCTGCAATGAA 376  
 4622 TTTTGGGTAATTTTATTTGTTATTTATTTATTTTGTATTTTGTATTTTGTAGTTT 4681  
 377 TAGGTATGATCTAAATCTTGTAAATTTGTTG 408  
 4682 TATGGGATTAATTTTGTGTTTATGTTTATG 4713

RESULT 7  
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 LOCUS AX344561 349980 bp DNA linear PAT 01-FEB-2002  
 DEFINITION Sequence 12 from Patent WO0200932.  
 ACCESSION AX344561  
 VERSION AX344561.1 GI:18492447  
 KEYWORDS synthetic construct  
 SOURCE synthetic construct  
 ORGANISM artificial sequences.

REFERENCE  
 1  
 Olek.A., Piepenbrock,C. and Berlin,K.  
 TITLE Diagnosis of known genetic parameters within the mhc  
 JOURNAL Patent: WO 0200932-A 12 03-JAN-2002;  
 EpiGenomics AG (DE)  
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 Location/Qualifiers

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ORIGIN  
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 Best Local Similarity 47.8%; Pred. No. 0.091;  
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 QY 98 GCGAGATTGTTCTATCTTGGCCATTAATCTTCTTTCGCGAATGATCTTCCGTAT 157  
 Db 277783 TTTAGTAAATTTTATTTTTCGATGTTTTTATTTTATTTTTCGAGTTTAAATAT 277842  
 QY 158 AAGATTTTAGTTAGAGATGAATCGTATAGCTAGATTTTATCACCAGATGTTCTTGT 217  
 Db 277843 ATATATTATTAATAATTTGGAGTTTTTATTTTATTTATATAGATAAGTTTTTATTA 277902  
 QY 218 TCTAGATCTCTGAATCTCGATAGTTTTCACATGTTGAATAGATTTCTTATTCGG 277  
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 QY 278 CGATTGTTGATTAGGTTTTGATTTTCTTGTATGCGATTGCAATTAGGATTTCTTT 337  
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RESULT 8  
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 DEFINITION Plasmodium falciparum 3D7 chromosome 11 section 7 of 8 of the complete sequence.  
 ACCESSION AE014842 AE014186  
 VERSION AE014842.1 GI:23496321  
 KEYWORDS Plasmodium falciparum 3D7  
 SOURCE Plasmodium falciparum 3D7  
 ORGANISM Plasmodium falciparum 3D7  
 REFERENCE 1 (bases 1 to 253151)  
 AUTHORS Gardner,M.J., Hall,N., Fung,E., White,O., Berriman,M., Hyman,R.W., James,K., Eissen,J.A., Rutherford,K., Salzberg,S.L., Craig,A., Kyes,S., Chan,M.-S., Nene,V., Shallow,S.J., Sub,B., Peterson,J., Anguoli,S., Percey,M., Allen,J., Selengut,J., Haft,D., Mather,M.W., Vaidya,A.B., Martin,D.M.A., Fairlamb,A.H., Fraunholz,M.J., Roos,D.S., Ralph,S.A., McFadden,G.I., Cummings,L.M., Subramanian,G.M., Mungall,C., Venter,J.C., Carucci,D.J., Hoffman,S.L., Newbold,C., Davis,R.W., Fraser,C.M. and



```

Barrell,B.
Genome sequence of the human malaria parasite Plasmodium falciparum
JOURNAL Nature 419 (6906), 498-511 (2002)
PUBMED 12368864
REFERENCE 2 (bases 1 to 253151)
AUTHORS Gardner,M.J.
TITLE Direct Submission
JOURNAL Submitted (13-SEP-2002) The Institute for Genomic Research, 9712
Medical Center Dr., Rockville, MD 20850, USA
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pieces.
ACCESSION
BX470257
KEYWORDS
HTG; HTGS PHASE1; HTGS DRAFT; HTGS_FULLTOP.
SOURCE
Danio rerio (zebrafish)
ORGANISM
Danio rerio
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;
Cypriniformes; Cyprinidae; Danio.
1 (bases 1 to 206653)
REFERENCE
1 McLaren,S.
DIRECT SUBMISSION
Submitted (24-SEP-2003) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
zfish-help@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
On Sep 24, 2003 this sequence version replaced gi:32169046.
COMMENT
----- Genome Center
Center: Wellcome Trust Sanger Institute
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: zfish-help@sanger.ac.uk
----- Project Information
Center project name: zK99015
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Chemistry: Dye-terminator; 100% of reads
Consensus quality: 205905 bases at least Q40
Consensus quality: 206063 bases at least Q30
Consensus quality: 206151 bases at least Q20
Insert size: 206253; sum-of-contigs
Quality coverage: 7.80x in Q20 bases; sum-of-contigs Quality
coverage: 8.14x in Q20 bases; agarose-fp
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 5 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 64965. contig of 64965 bp in length

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## artificial sequences.

## REFERENCE

1  
Olek, A., Piepenbrock, C. and Berlin, K.  
Diagnosis of known genetic parameters within the mhc  
Patent: WO 0200932-A 24 03-JAN-2002;  
Epigenomics AG (DE)

## FEATURES

## source

Location/Qualifiers  
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## ORIGIN

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Best Local Similarity 45.2%; Fred. No. 0.37; Indels 0; Gaps 0;  
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Db 317872 AATTAATTTTGTAGTTTGTAGTAAATTTTTCGATTTTTTTTTTTTTTTAGATTTTT 317931  
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Db 318172 TTTTGTATTTTAAATAATTTTTTTTTTTTATTTTTTTT 318209

Search completed: June 20, 2004, 03:45:45  
Job time : 2934.58 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: June 19, 2004, 23:55:13 ; Search time 309.833 Seconds

(without alignments)

5731.297 Million cell updates/sec

Title: US-09-000-062-6

Perfect score: 418

Sequence: 1 tggagtaacattcttcgac.....taattgtgacagatccc 418

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 337863 seqs, 2124099041 residues

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N Geneseq 29Jan04:\*

1: Geneseqn1980s:\*

2: Geneseqn1990s:\*

3: Geneseqn2000s:\*

4: Geneseqn2001as:\*

5: Geneseqn2001bs:\*

6: Geneseqn2002s:\*

7: Geneseqn2003as:\*

8: Geneseqn2003bs:\*

9: Geneseqn2003cs:\*

10: Geneseqn2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	418	100.0	418	2	Aat85996 Arabidops
2	54.4	13.0	598	6	Ab133634 Human inm
3	50.6	12.1	17527	6	Ab133432 Human inm
4	50.6	12.1	17527	6	Aas63332 Chemicall
5	50.2	12.0	6306	4	Aas45515 Chemicall
6	50.2	12.0	6306	4	Abk28457 DNA trans
7	49	11.7	5987	6	Ab133562 Human inm
8	49	11.7	5987	6	Abg67101 Human ang
9	49	11.7	5987	9	Abd54307 Pretreate
10	49	11.7	5987	9	Abd54179 Pretreate
11	48.8	11.7	15479	6	Abk39964 Human che
12	48.4	11.6	17848	4	Aas45323 Chemicall
13	48.4	11.6	17848	6	Abk39976 Human che
14	48.4	11.6	17848	6	Abk28164 DNA trans
15	48	11.5	2501	7	Abz10074 Haematopo
16	48	11.5	2501	7	Abz10220 Haematopo
17	48	11.5	6059	4	Ab133480 Human inm
18	48	11.5	8245	4	Aas46448 Tumour su
19	48	11.5	16724	6	Ab133090 Human inm
20	48	11.5	16724	6	Ab134536 Human met
21	48	11.5	16724	6	Ab170259 Chemicall
22	47.6	11.4	5649	9	Abd54254 Pretreate
23	47.6	11.4	5649	9	Ade84184 Human lym

24	47.4	11.3	21537	6	Ab133999 Human inm
25	47.2	11.3	5371	6	Ab134295 Human inm
26	46.8	11.2	714	6	Abq18144 Oligonuel
27	46.8	11.2	714	6	Abq18145 Oligonuel
28	46.8	11.2	8805	6	Abk40016 Human che
29	46.6	11.1	6012	6	Ab134564 Human met
30	46.6	11.1	6012	6	Ab170289 Chemicall
31	46.6	11.1	6668	6	Ab133697 Human inm
32	46.2	11.1	6048	6	Ab132509 Human inm
33	46.2	11.1	18817	6	Ab134495 Human met
34	46.2	11.1	18817	6	Ab170162 Chemicall
35	46.2	11.1	26811	2	Aax20253 Borrelia
36	46	11.0	5649	4	Aas46384 Tumour su
37	46	11.0	5649	6	Abk40008 Human che
38	46	11.0	5649	6	Ab132849 Human inm
39	46	11.0	5649	9	Abd54126 Pretreate
40	46	11.0	5649	9	Ade84108 Human lym
41	46	11.0	6161	6	Ab132623 Human inm
42	46	11.0	7892	6	Abk40056 Human che
43	45.8	11.0	6223	6	Aas61176 Human gen
44	45.6	10.9	374	7	Abx47736 Bovine ES
45	45.6	10.9	593	6	Abq20161 Oligonuel

ALIGNMENTS

RESULT 1

AAT85996

ID AAT85996 standard; DNA; 418 BP.

XX

AC AAT85996;

XX

DT 17-NOV-1997 (first entry)

XX

DE Arabidopsis thaliana histone H3.3-like DNA fragment (intron 1).

XX

KW Plant expression regulation sequence; intron 1; histone;

KW herbicide tolerance; 5-enolpyruvylshikimate-3-phosphate synthase; EPSPS;

KW glyphosate; ds.

XX

OS Arabidopsis thaliana.

XX

PH Key

intron

FT Location/Qualifiers

FT 1..418

FT /\*tag= a

FT /number= 1

XX

PN WO9704114-A2.

XX

PD 06-FEB-1997.

XX

PF 17-JUL-1996; 96WO-FR001109.

XX

PR 19-JUL-1995; 95FR-00008980.

XX

PA (RHON ) RHONE POULENC AGROCHIMIE.

XX

PI Derose R, Chaubet N, Gigot C;

XX

DR WPI; 1997-132652/12.

XX

PT New regulatory sequence for chimeric gene expression in rapidly growing

PT parts of a plant - includes at least one intron from a plant histone gene

PT and is useful for imparting resistance to herbicides.

XX

PS Claim 4; Page 26; 31pp; French.

XX

CC The known cosmid clone c22 of Arabidopsis thaliana contains two histone

CC H3.3-like genes. Digestion of clone c22 with restriction enzyme DdeI,

CC Klenow fragment and MspI generated a fragment of 418 bp having the

CC present sequence. This fragment, designated intron 1, was ligated to

CC synthetic linkers for cloning into plant expression vectors. In addition

CC

CC to the intron 1 sequence, the vectors contained a plant promoter and a  
 CC herbicide tolerance gene (e.g. a mutated version of the 5'  
 CC enolpyruvylshikimate-3-phosphate synthase (EPSPS) gene). The intron  
 CC enhances expression of the herbicide tolerance gene in rapidly growing  
 CC parts of plants. The intron can also be used to enhance expression of  
 CC genes that impart resistance to pathogens or that encode nutritional or  
 CC therapeutic proteins  
 XX Sequence 418 BP; 100 A; 63 C; 85 G; 170 T; 0 U; 0 Other;  
 SQ

Query Match 100.0%; Score 418; DB 2; Length 418;  
 Best Local Similarity 100.0%; Pred. No. 2.8e-92;  
 Matches 418; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGAGGTAGATCTTCGATCTCTTGGATTTCTCGTGAATATTTTCGTGATCTGGA 60  
 DB 1 TGAGGTAGATCTTCGATCTCTTGGATTTCTCGTGAATATTTTCGTGATCTGGA 60

QY 61 AACTACTCGAATCGCTCGATAGGTGGTACGAAATTAGCGGAGATTAGTTCTATCTTGG 120  
 DB 61 AACTACTCGAATCGCTCGATAGGTGGTACGAAATTAGCGGAGATTAGTTCTATCTTGG 120

QY 121 CAATATCTTGTCTTTCGCGGATGATCTCCGTATAAAGATTTAGGTTAGAGATGAA 180  
 DB 121 CAATATCTTGTCTTTCGCGGATGATCTCCGTATAAAGATTTAGGTTAGAGATGAA 180

QY 181 TCGTATAGCTAGATTTTCATCACCAGATAGTTCTTTGTCTAGAAATCTCGAAATTCGGA 240  
 DB 181 TCGTATAGCTAGATTTTCATCACCAGATAGTTCTTTGTCTAGAAATCTCGAAATTCGGA 240

QY 241 TAGTTTTCACATGCTGTAATAGATTTCTTATTCGGCGATGTTGATTAGGTTTGGAT 300  
 DB 241 TAGTTTTCACATGCTGTAATAGATTTCTTATTCGGCGATGTTGATTAGGTTTGGAT 300

QY 301 TTTCCTGATTATGCGATTCGAAATAGGATTTCTTTGGTTTGTGATCTTACGATA 360  
 DB 301 TTTCCTGATTATGCGATTCGAAATAGGATTTCTTTGGTTTGTGATCTTACGATA 360

QY 361 CATTCTCGAATGAAATAGATGATGATCTAAATCTTGTGTTGTTGAAACAGATCCC 418  
 DB 361 CATTCTCGAATGAAATAGATGATGATCTAAATCTTGTGTTGTTGAAACAGATCCC 418

RESULT 2  
 ABL33634  
 ID ABL33634 standard; DNA; 5998 BP.  
 XX  
 AC ABL33634;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Human immune system associated gene SEQ ID NO: 1607.  
 XX  
 KW Human; immune system disease; cytosine methylation; antiasthmatic;  
 KW antiarteriosclerotic; antianaemic; cytosine methylation; antiasthmatic;  
 KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;  
 KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;  
 KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;  
 KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
 KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
 ds.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200200928-A2.  
 XX  
 XX  
 PD 03-JAN-2002.  
 XX  
 XX  
 PF 02-JUL-2001; 2001WO-EP007537.  
 XX  
 XX  
 PR 30-JUN-2000; 2000DB-01032529.  
 PR  
 PR 01-SEP-2000; 2000DB-01043826.  
 XX

PA (EPIG-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K;  
 XX  
 DR WPI; 2002-130909/17.  
 XX  
 PT Nucleic acid comprising fragment of chemically modified gene, useful for  
 PT diagnosis and treatment of diseases associated with abnormal cytosine  
 PT methylation.  
 XX  
 PS Claim 1; SEQ ID NO 1607; 32pp + Sequence Listing; German.  
 XX  
 CC The present invention provides a number of human immune system associated  
 CC genes which are modified by the methylation of cytosines. The sequences  
 CC can be used in the diagnosis and treatment of immune system disorders,  
 CC including eye diseases such as retinopathy, neovascular glaucoma and  
 CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
 CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
 CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
 CC diseases. The present sequence is a gene of the invention  
 XX  
 SQ Sequence 5998 BP; 1217 A; 92 C; 1806 G; 2883 T; 0 U; 0 Other;  
 Query Match 13.0%; Score 54.4; DB 6; Length 5998;  
 Best Local Similarity 49.7%; Pred. No. 0.0013;  
 Matches 165; Conservative 0; Mismatches 166; Indels 1; Gaps 1;

QY 78 GATGATGTTGATGAAATAGCGGAGATTAGTTCTTATCTCGCCATTAATCTGTTCTT 137  
 DB 4382 GACTAGTGGTTGAGTATAGTTAAATGTTAGTTTCGAATTTTGGGTTAAATGATTTT 4441

QY 138 CGCGGATGATCTTCGGTA-TAAGATTTTAGTTAGATGAATCGTATAGATTT 196  
 DB 4442 TGTTTAGTTTGTGATAGTTAGTTGGGATATAGTTATAGTTTGTGTTAATTA 4501

QY 197 CATCACCATAGTTTCTTTTGTCTAGATCTCTGAAATCTCGATAGTTTTCACATGCT 256  
 DB 4502 AAAAAAATGTTTTTTTTTTTATAGAGATAGAAGTTTTTTTGTGTTAGTTGG 4561

QY 257 AATAGATGTTCTTATTCGGGATGTTGATAGGTTTGTGATTTCTGTTGATGCA 316  
 DB 4562 TTTTGAATTTTGGTTTGGGATTTTATTTTTTTTGTGTTTGTGTTATATA 4621

QY 317 TTGCAATTAGGGATTTTCTTTTGTGTTGATCTTACGATACATCTCTGCAATGAA 376  
 DB 4622 TTTTAGGTAATTTTATTTGTTATTTATTTATTTTGTGTTTGTGTTT 4681

QY 377 TACGTATGGAATCTAAATCTTGTGTTAATTTGTTG 408  
 DB 4682 TATGGGATTTTGTGTTTGTGTTTATGTTTGTAG 4713

RESULT 3  
 ABL33432  
 ID ABL33432 standard; DNA; 17527 BP.  
 XX  
 AC ABL33432;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Human immune system associated gene SEQ ID NO: 1405.  
 XX  
 KW Human; immune system disease; cytosine methylation; antiasthmatic;  
 KW antiarteriosclerotic; antianaemic; cytosine methylation; antiasthmatic;  
 KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;  
 KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;  
 KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;  
 KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
 KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
 ds.  
 XX  
 OS Homo sapiens.  
 XX



PN	WO200200928-A2.
XX	
PD	03-JAN-2002.
XX	
PF	02-JUL-2001; 2001WO-EP007537.
XX	
PR	30-JUN-2000; 2000DE-01032529.
PR	01-SEP-2000; 2000DE-01043826.
XX	(EPIG-) EPIGENOMICS AG.
PA	
PI	Olek A, Piepenbrock C, Berlin K;
XX	WPI; 2002-130909/17.
XX	
PT	Nucleic acid comprising fragment of chemically modified gene, useful for
PT	diagnosis and treatment of diseases associated with abnormal cytosine
PT	methylation.
XX	
PS	Claim 1; SEQ ID NO 1405; 32pp + Sequence Listing; German.
XX	
CC	The present invention provides a number of human immune system associated
CC	genes which are modified by the methylation of cytosines. The sequences
CC	can be used in the diagnosis and treatment of immune system disorders,
CC	including eye diseases such as retinopathy, neovascular glaucoma and
CC	macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
CC	leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
CC	rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
CC	diseases. The present sequence is a gene of the invention
XX	
SQ	Sequence 17527 BP; 4501 A; 296 C; 4094 G; 8636 T; 0 U; 0 Other;
	Query Match 12.1%; Score 50.6; DB 6; Length 17527;
	Best Local Similarity 52.1%; Pred. No. 0.013;
	Matches 113; Conservative 0; Mismatches 104; Indels 0; Gaps 0;
QY	192 GATTTCATCACGAGTAGTTCTTCTGTCTAGAAATCTCGAAATCTCGATAGTTTTCA 251
Db	16590 GTTGTTATAAGAATAATTTGGTTTTTATTGAGTCGTTATTATTAGAACGTTTTAAGA 16649
QY	252 TGCGTAATAAGATTGTTCTATTCCGCCGATTGTTGATTAGGCTTTGATTTCTTGATTA 311
Db	16650 GGTCGGTATTGTTGGTTGTTTTTTTGGTTTTTTTTTTTAAATTTTTTTTAGTTT 16709
QY	312 TGCGATTGCAAATAGGGATTTCTTTGGTTTTGTGTGATCTACGATACATTCCTCGAA 371
Db	16710 TAATAGTGATTATGGGAAGTTTTTTTTTTGTTTTGGTTGTTTTTGTATTTTTTAT 16769
QY	372 TTGAATACGTATCGATCATAAATCTGTTAATTGTTG 408
Db	16770 TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTAG 16806
RESULT 4	
AAS63332	
ID	AAS63332 standard; DNA; 17527 BP.
XX	
AC	AAS63332;
XX	
DT	29-JAN-2002 (first entry)
XX	
DE	Chemically pretreated metabolism associated gene #27.
XX	
KW	Human; cytostatic; anti-tumour; metabolism; metabolic disease; liver;
KW	solid tumour; cancer; cytosine methylation; epigenetic; eye; kidney;
KW	single nucleotide polymorphism detection; SNP; stool; urine; lung;
KW	cerebral-spinal fluid; intestine; brain; heart; prostate; breast; DUSP2;
KW	EPHX2; QPBR; SGSH; SHMT2; SLC7A2; SLC7A4; TYMS; ds.
XX	
OS	Homo sapiens.
XX	
PX	WO200176451-A2.
FN	

18-OCT-2001.  
06-APR-2001; 2001WO-EP004016.  
06-APR-2000; 2000DE-01019058.  
07-APR-2000; 2000DE-01019173.  
30-JUN-2000; 2000DE-01032529.  
01-SEP-2000; 2000DE-01043826.  
(EPIG-) EPIGENOMICS AG.  
Olek A, Piepenbrock C, Berlin K;  
WPI; 2002-010834/01.  
New nucleic acid, useful for diagnosis and therapy of metabolic disease, solid tumor and cancers, comprises segment of chemically modified genomic sequences of genes associated with metabolism.  
Claim 1; Page 79-84; 143pp; English.  
The invention relates to a nucleic acid (I) comprising a sequence at least 18 bases of a segment of the chemically pretreated DNA of genes associated with metabolism such as DUSP2 (NM\_004418), EPHX2 (NM\_001979), QPDR (NM\_000320), SGSH (NM\_000199), SHMT2 (NM\_005412), SLC7A2 (NM\_003046), SLC7A4 (NM\_004173) and TMS (NM\_001071) (all undefined). (I) are useful for diagnosis and therapy of metabolic disease, solid tumours and cancers; as primer oligonucleotides for the amplification of DNA sequences, for detecting the cytosine methylation state and/or single nucleotide polymorphisms (SNPs) in a chemically treated DNA of genes associated with metabolism. An array of (I) is useful for ascertaining genetic and/or epigenetic parameters for the diagnosis and/or therapy of existing diseases or the predisposition to specific diseases by analysing cytosine methylations. The method involves chemically treating genomic DNA sample by a solution of bisulphite, hydrogen sulphite or disulphite such that cytosine bases which are unmethylated at the 5th-position are converted to uracil or another base which is dissimilar to cytosine in terms of hybridisation behaviour and amplifying fragments of the chemically pretreated genomic DNA. The genomic DNA is from cells or cellular components which contain DNA, sources of DNA comprising, for e.g. cell lines, biopsies, blood, sputum, stool, urine, cerebral-spinal fluid, tissue embedded in paraffin such as tissue from eye, intestine, kidney, brain, heart, prostate, lung, breast or liver, histologic object slides and their combinations. Genetic parameters are mutations, in particular insertions, deletions, point mutations, inversions and polymorphisms of genes associated with metabolism and sequences further required for their regulation. Epigenetic parameters are in particular cytosine methylations and further chemical modifications of DNA bases of genes associated with metabolism. Further epigenetic parameters include for e.g. the acetylation of histones which correlates with DNA methylation. AAS63306-AAS63373 represent chemically pretreated metabolism associated genes, and related primers of the invention

XX	SQ	Sequence	17527 BP; 4501 A; 296 C; 4094 G; 8636 T; 0 U; 0 Other;
	Query Match	12.1%; Score 50.6; DB 6; Length 17527;	
	Best Local Similarity	52.1%; Pred. No. 0.013;	
	Matches	113; Conservative 0; Mismatches 104; Indels 0; Gaps 0;	
QY	192	GATTTCATCACCAGATAGTTCTTTGTCFAGAAATCTCGAATAGTTTTCA	251
Db	16590	GTTCGTTAAGAATAATTTGTTTTTATTGAGTCGTTATTATTAGAAAGTTTTA	16649
QY	252	TGTTGTAATAGATGTCTTATTCGGCGATGTTCGATTAGGGTTTGATTTCTTGATTA	311
Db	16650	GGTGGATTTGTGGGTGTTTTTGTGGTTTTTTTTTTTATTTTTTTTTTTAGTTTT	16709
QY	312	TGCGATTGCAATPAGGATTTCTTTGGSTTTTGTTGATCTTACGATACATTTCTCTCGAA	371
Db	16710	TAATAGTGTATTATGGGAAGTTTTTTTTTGTTTTGTGTTTTTTTGATTTTTTTAT	16769
QY	372	TTGAATACGTATCGATCTAAATCTCTGTTAAATTTGTTG	408

	SQ	Sequence	17527 BP; 4501 A; 296 C; 4094 G; 8636 T; 0 U; 0 Other;
RESULT 4			
AAS63332		Query Match	12.1%; Score 50.6; DB 6; Length 17527;
ID AAS63332 standard; DNA; 17527 BP.		Best Local Similarity	52.1%; Pred No. 0.013;
XX AC		Matches 113; Conservative	0; Mismatches 104; Indels 0; Gaps 0;
AAS63332;			
XX XX			
DT 29-JAN-2002 (first entry)	QY	192 GAATTCATCACAGATAGTTCTTGTCTGATGAATCTCTCAAAATTCTCGATAGTTTTACAA	251
Chemically pretreated metabolism associated gene #27.	Db	16590 GTTGTTATAAGAATAATTTGGTTTTTATTTAGTCGTTATTATTATTAAGAAGTTTAAAGA	16649
Human; cytostatic; anti-tumour; metabolism; metabolic disease; liver;	QY	252 TGCTAATAATAGATTGTTCTTATTTCGCCGATTCTGCATTAGGTTTTGATTTCTTGATTA	311
solid tumour; cancer; cytosine methylation; epigenetic; eye; kidney;	Db	16650 GGTCGATTATGTTGGTGTGTTTTTTGTTGGTTTTTTTTTTATTTTTTTTAGTTT	16709
single nucleotide polymorphism detection; SNP; stool; urine; lung;	QY	312 TGCATATGCATATAGGAGATTTCTTGTTGGTTTTGCTGTCATCTTACGATACAATCCTCGAA	371
cerebral-spinal fluid; intestine; brain; heart; prostate; breast; DUSP2;	Db	16710 TAATAGTGAATATGGGAAGTTTTTTTTTGTTTTTGTTGTTTTTGTATTTTTTAT	16769
EPHX2; QDPR; SGSH; SMT2; SLC7A2; SLC7A4; TYMS; ds.	QY	372 TTGAATACGATATGATCTAAATCTTCTTAATTTGTTG	408
Homo sapiens.			
WO200176451-A2.			
XX PN			
PX PX			

Db		16770 TTTTTCGTTATGTTTTTTAGATTTTGTGGTTTATTGTTTATTTTAAAGTT 5609	
	RESULT 5		
	AAS45515		
ID	AAS45515 standard; DNA; 6306 BP.		
XX	AC		
XX	AC		
XX	XT		
DT	18-DEC-2001 (first entry)		
XX			
DE	Chemically pretreated complementary DNA associated with cell cycle #110.		
XX			
KW	Cell cycle; human; CpG dinucleotide; cytosine methylation; HIV; aging;		
KW	human immunodeficiency virus; neurodegenerative disorder; solid tumour;		
KW	graft-versus-host disease; glomerular disease; Lewy body disease; cancer;		
KW	arthritis; arteriosclerosis; anti-HIV, neuroprotective; antiarthritic;		
KW	immunosuppressive; antitumour; cytostatic; antiarteriosclerotic; ds;		
KW	PCR primer.		
OS	Homo sapiens.		
PX			
PN	WO200168911-A2.		
XX			
PD	20-SEP-2001.		
XX			
PF	15-MAR-2001; 2001WO-EF002945.		
XX			
PR	15-MAR-2000; 2000DE-01013847.		
PR	06-APR-2000; 2000DE-01019058.		
PR	07-APR-2000; 2000DE-01019173.		
PR	30-JUN-2000; 2000DE-01032529.		
PR	01-SEP-2000; 2000DE-01043826.		
XX			
PA	(EPIG-) EPIGENOMICS AG.		
PI	Olek A, Piepenbrock C, Berlin K;		
DR	WTI; 2001-602751-68.		
XX			
PT	Designing primers and probes for analyzing diseases associated with		
PT	cytosine methylation state e.g. arthritis, cancer, aging,		
PT	arteriosclerosis comprising fragments of chemically modified genes		
PT	associated with cell cycle.		
XX			
PS	Claim 1; SEQ ID NO 220; 28pp; English.		
XX			
CC	Sequences AAS45296-AAS45520 represent chemically pretreated genomic DNA		
CC	molecules associated with the cell cycle and specific PCR primers of the		
CC	invention. The sequences are useful for detecting the methylation state		
CC	of all CpG dinucleotides in a sequence and therefore for analysing		
CC	associated diseases. By analysing cytosine methylations in the pretreated		
CC	DNA, genetic and/or epigenetic parameters for the diagnosis and therapy		
CC	of existing diseases or the predisposition to specific diseases can be		
CC	ascertained. The parameters may be compared to another set of genetic		
CC	and/or epigenetic parameters, the differences serving as basis for		
CC	diagnosis and/or prognosis events which are disadvantageous to patients.		
CC	The sequences of the invention are useful for the diagnosis and therapy		
CC	of HIV infection, neurodegenerative disorders, graft-versus-host disease,		
CC	aging, glomerular disease, Lewy body disease, arthritis,		
CC	arteriosclerosis, solid tumours and cancers		
SQ	Sequence 6306 BP; 1455 A; 190 C; 1580 G; 3081 T; 0 U; 0 Other;		
	Query Match 12.0%; Score 50.2; DB 4; Length 6306;		
	Best Local Similarity 50.2%; Pred. No. 0.014;		
	Matches 124; Conservative 0; Mismatches 123; Indels 0; Gaps 0;		
Qy	161 GATTTTAGGTAGATGAATCGTATAGCTAGTATTCATCACCAGATAGTTCTTGTC 220		
Db	5490 GAGTTTCGGTTCGAGGGGTATGATGTTAGTATTAATTTCTTTTAGGTAG 5549		
	221 AGAATCTCTCAATCTCGCATGTTTTTCATCTGTAATAATAGATTTCTTTATCGGCA 280		

CC deficiency, viral infection, retroviral infection, Sezary syndrome,  
 CC haematological disorders, immunological disorders, Werner syndrome,  
 CC tuberculosis, developmental disorders, psoriasis, Rieger's syndrome,  
 CC neurological disorders, neurodegenerative disorders, Waardenburg  
 CC syndrome, Niemann-Pick disease, myelodysplastic syndrome, myocardial  
 CC infarction, hypertension, angiogenesis, erythropoiesis, congenital heart  
 CC disease, HDR syndrome, arthritis, polyglutamine disorders, solid tumours  
 CC or cancer. Sequences ABK28127-ABK28472 represent DNA transcription  
 CC associated genomic DNA molecules of the invention. Note: The sequence  
 CC data for this patent did not form part of the printed specification but  
 CC was obtained in electronic format directly from the European Patent  
 CC Office

XX Sequence 6306 BP; 1455 A; 190 C; 1580 G; 3081 T; 0 U; 0 Other;  
 SQ

Query Match 12.0%; Score 50.2; DB 6; Length 6306;  
 Best Local Similarity 50.2%; Pred. No. 0.014;  
 Matches 124; Conservative 0; Mismatches 123; Indels 0; Gaps 0;

QY 161 GATTGTTAGTTAGATGATCGTATAGCTAGATTCATCAGATAGTTCTTCT 220  
 |||||  
 DB 5490 GAGTTTCGGTTCAGGGGTATAGTATGTTAGTATGATTAATTTGTTTATAGGTAG 5549  
 |||||

QY 221 AGAATCTCTGMAATCTCGATAGTTTTCACATGCTGTAATAGATGTTCTTATTCGGCGA 280  
 |||||  
 DB 5550 AIGTTTTCGTATGATTTAGATTTTGTGTTTGTATTTTATTTTAAAGTT 5609  
 |||||

QY 281 TTGTTGATAGGTTTGTGATTTTCTGATATGCGATTCGATAGGATTTCTTGGT 340  
 |||||  
 DB 5610 TGATTTAAATTTTGTGTTGTTTATATATCGTTTGTGTTTGTGTTTGTGTTT 5669  
 |||||

QY 341 TTGTTGTTGATCTTACGATACATTCCTGCAATTTGAAATAGTATGGAATCTGTTA 400  
 |||||  
 DB 5670 TTTTGTGTTGATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 5729  
 |||||

QY 401 ATTGTT 407  
 |||||  
 DB 5730 TTTTGT 5736  
 |||||

RESULT 7  
 ABL33562  
 ID ABL33562 standard; DNA; 5987 BP.  
 XX  
 AC ABL33562;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Human immune system associated gene SEQ ID NO: 1535.  
 XX  
 KW Human; immune system disease; cytosine methylation; antiasthmatic;  
 KW antiarteriosclerotic; antianaemic; cytostatic; nootropic;  
 KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;  
 KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;  
 KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;  
 KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
 KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
 KW ds.  
 XX  
 OS Homo sapiens.  
 XX  
 EN WO200200928-A2.  
 XX  
 PD 03-JAN-2002.  
 XX  
 PF 02-JUL-2001; 2001WO-EP007537.  
 XX  
 PR 30-JUN-2000; 2000DE-01032529.  
 PR 01-SEP-2000; 2000DE-01043826.  
 XX  
 FA (EPIG-) EPIGENOMICS AG.  
 XX  
 PI Olek A, Piepenbrock C, Berlin K;

XX WPI; 2002-130909/17.  
 XX Nucleic acid comprising fragment of chemically modified gene, useful for  
 XX diagnosis and treatment of diseases associated with abnormal cytosine  
 XX methylation.  
 XX Claim 1; SEQ ID NO 1535; 32pp + Sequence Listing; German.  
 XX The present invention provides a number of human immune system associated  
 XX genes which are modified by the methylation of cytosines. The sequences  
 XX can be used in the diagnosis and treatment of immune system disorders,  
 XX including eye diseases such as retinopathy, neovascular glaucoma and  
 XX macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
 XX leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
 XX rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
 XX diseases. The present sequence is a gene of the invention  
 XX

SQ Sequence 5987 BP; 1754 A; 96 C; 1096 G; 3041 T; 0 U; 0 Other;  
 Query Match 11.7%; Score 49; DB 6; Length 5987;  
 Best Local Similarity 47.5%; Pred. No. 0.026;  
 Matches 145; Conservative 0; Mismatches 160; Indels 0; Gaps 0;

QY 108 TTTCTATTCTGGCCATTATCTTGTCTTCGCGGAATGATCTTCGATATAAGATTTTA 167  
 |||||  
 DB 2953 TTTTGTGTTAGTTAGATTTGTTATTTTAAATTTTGTATTTTAAAGTTTA 2912  
 |||||

QY 168 GGTAGATGATCGTATAGCTAGATTCATCAGATAGTTCTTCTCTAGATCT 227  
 |||||  
 DB 2913 GATTTTGTGAAATTTTGTGTTTGTATTTTAAATTTTGTGAAATTTTATTTTATTT 2972  
 |||||

QY 228 CTGAAATCTCGATAGTTTTCACATGCTGTAATAGATTTCTTATTCGCGGATTTGTA 287  
 |||||  
 DB 2973 GTTATAAGTGATAGATTTTATTTATGGAATGATGGAGTTAAATTTTGTAGTTTA 3032  
 |||||

QY 288 TTAGGTTTGTGATTTCTGATATGCGATTCGAAATAGGATTTCTTGGTTTCTGT 347  
 |||||  
 DB 3033 AAATAATTTAAATATAATTTTATTTTGTGTTTGTGTTTGTGTTTGTGTTTAAAT 3092  
 |||||

QY 348 TGATCTTACGATACATTCCTGCAATTCGAATGATGATCTAAATCTTGTAAATTTGTT 407  
 |||||  
 DB 3093 TTTTGTGATATAGATTAATTAATGTTTGTATAGGATAGATGATGATGATTTT 3152  
 |||||

QY 408 GAACA 412  
 |||||  
 DB 3153 TAAAA 3157  
 |||||

RESULT 8  
 ABL33562  
 ID ABL33562 standard; DNA; 5987 BP.  
 XX  
 AC ABL33562;  
 XX  
 DT 28-AUG-2002 (first entry)  
 XX  
 DE Human angiogenesis associated polynucleotide SEQ ID NO 131.  
 XX  
 KW Human; angiogenesis; methylation; eye disease; glaucoma; tumour;  
 KW inflammation; rheumatoid arthritis; diabetic retinopathy; antileucers;  
 KW macular degeneration; inflammatory bowel disease; Crohn's disease;  
 KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;  
 KW antiarteriosclerotic; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 EN WO2002046454-A2.  
 XX  
 PD 13-JUN-2002.  
 XX  
 PF 06-DEC-2001; 2001WO-EP014320.  
 XX

PR 06-DEC-2000; 2000DE-01061338.  
XX (EPIG-) EPIGENOMICS AG.  
XX Schacht O;  
XX WPI; 2002-500450/53.  
XX New nucleic acid fragments from chemically treated angiogenesis-  
PT associated genes, useful for determining methylation status, e.g. in  
PT diagnosis or treatment of cancer.  
XX Claim 1; SEQ ID NO 131; 41pp + Sequence Listing; German.  
XX  
XX The invention relates to a nucleic acid (I) comprising a segment of 18  
CC bases of chemically pretreated DNA of angiogenesis-associated genes (II)  
CC having sequences (ABQ66971-ABQ67178) or their complements. (I), also  
CC related oligomers, are used to evaluate the methylation status and/or  
CC single-nucleotide polymorphisms, in angiogenesis-related genes, for  
CC diagnosis and treatment of eye diseases, proliferative retinopathy,  
CC neovascular glaucoma, solid tumors, inflammation, rheumatoid arthritis,  
CC diabetic retinopathy, macular degeneration caused by neovascularisation,  
CC psoriasis, arteriosclerosis, inflammatory bowel diseases, ulcers and  
CC Crohn's disease. Note: The sequence data for this patent did not form  
CC part of the printed specification, but was obtained in electronic format  
CC directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences  
XX  
SQ Sequence 5987 BP; 1754 A; 96 C; 1096 G; 3041 T; 0 U; 0 Other;  
Query Match 11.7%; Score 49; DB 6; Length 5987;  
Best Local Similarity 47.5%; Pred. No. 0.026;  
Matches 145; Conservative 0; Mismatches 160; Indels 0; Gaps 0;  
QY 108 TTCTATCTTCGCCATATCTTGTCTTCGCCGAATGATCTCCGATATAAGATTTA 167  
DB 2853 TTTTCTTTTATGTAGATTGTAATTTTTTTTAAATTTTCTTTTATTTTAAAGTTTA 2912  
QY 168 GGTAGAGATGAATCGTAGCTAGATTTTCATCCAGATAGTTCTTTGCTAGAACT 227  
DB 2913 GATTTTTTTGAAATTTTTTTTTTGAATTTTTTAAAGTATATGGAATTTTTTATTTATTTT 2972  
QY 228 CTGAATTCCTGATAGTTTTCACATGCTGTAATAGATTTCTTATTCGCCGATGTTGA 287  
DB 2973 GTTATATAAGTAGATAGATTTTATTTATGCGGATAGATGAGTTTAAATTTTTTGAGTTTA 3032  
QY 288 TTAGGTTTGTGATTTCTTGTGATTCGATTCGAATTCGGAATTTCTTTGTTTGTGT 347  
DB 3033 AAATAATTTAAATATAATTTATTTTATGTTTGTGTTTTTTTATTTTGTATTTAAAT 3092  
QY 348 TGATCTTACGATACATCTCCGCAATTCGATGATGATGATCTAAATCTGTTAAATTTGT 407  
DB 3093 TTTTCTTTAGATAATAGATAATTAATGTTTGTATAGGAAGATAATGATGATGATTTT 3152  
QY 408 GAACA 412  
DB 3153 TAAAA 3157  
RESULT 9  
ID ADB54307  
XX ADB54307 standard; DNA; 5987 BP.  
AC ADB54307;  
XX  
XX 04-DEC-2003 (first entry)  
XX Pretreated genomic DNA region 231.  
XX colon cell proliferative disorder; non methylated CpG dinucleotide;  
KW cytosstatic; cancer; adenoma; carcinoma; cytosine methylation state; ds.  
XX Unidentified.  
OS  
XX

PN WO2003072821-A2.  
XX  
XX 04-SEP-2003.  
XX 27-FEB-2003; 2003WO-EP002035.  
XX 27-FEB-2002; 2002EP-00004551.  
XX (EPIG-) EPIGENOMICS AG.  
XX Adorjan P, Burger M, Maier S, Nimmrich I, Becker E, Lesche R;  
PI Rujan T, Schmitt A;  
XX WPI; 2003-731620/59.  
XX  
XX Detecting and differentiating between colon cell proliferative disorders  
PT associated with a gene or its regulatory regions comprises contacting a  
PT target nucleic acid in a biological sample obtained from the subject with  
PT a reagent.  
XX  
XX Claim 32; SEQ ID NO 363; 74pp; English.  
XX  
XX The invention relates to a novel method for detecting and differentiating  
CC between colon cell proliferative disorders associated with at least one  
CC gene or its regulatory regions. The method comprises contacting a target  
CC nucleic acid in a biological sample obtained from the subject with at  
CC least one reagent or a series of reagents, where the reagent or series of  
CC reagents distinguishes between methylated and non methylated CpG  
CC dinucleotides within the target nucleic acid. The molecules of the  
CC invention demonstrate cytosstatic activity whilst the method may useful  
CC for detecting and differentiating between colon cell proliferative  
CC disorders, including cancers such as colon adenoma and colon carcinoma.  
CC The PNA (peptide nucleic acid)-oligomers are useful as probes for  
CC determining cytosine methylation state or single nucleotide  
CC polymorphisms. The current sequence is that of the pretreated genomic DNA  
CC region of the invention. This sequence is not shown within the  
CC specification but is taken from Wipoweb.  
XX  
SQ Sequence 5987 BP; 1754 A; 0 C; 1096 G; 3137 T; 0 U; 0 Other;  
Query Match 11.7%; Score 49; DB 9; Length 5987;  
Best Local Similarity 47.5%; Pred. No. 0.026;  
Matches 145; Conservative 0; Mismatches 160; Indels 0; Gaps 0;  
QY 108 TTCTATCTTCGCCATATCTTGTCTTCGCCGAATGATCTCCGATATAAGATTTA 167  
DB 2853 TTTTCTTTTATGTAGATTGTAATTTTTTTTAAATTTTCTTTTATTTTAAAGTTTA 2912  
QY 168 GGTAGAGATGAATCGTAGCTAGATTTTCATCCAGATAGTTCTTTGCTAGAACT 227  
DB 2913 GATTTTTTTGAAATTTTTTTTTTGAATTTTTTAAAGTATATGGAATTTTTTATTTATTTT 2972  
QY 228 CTGAATTCCTGATAGTTTTCACATGCTGTAATAGATTTCTTATTCGCCGATGTTGA 287  
DB 2973 GTTATATAAGTAGATAGAAATTTTATTTATGCGGATAGATGAGTTTAAATTTTTTGAGTTTA 3032  
QY 288 TTAGGTTTGTGATTTCTTGTGATTCGATTCGAATTCGGAATTTCTTTGTTTGTGT 347  
DB 3033 AAATAATTTAAATATAATTTATTTTATGTTTGTGTTTTTTTATTTTGTATTTAAAT 3092  
QY 348 TGATCTTACGATACATCTCCGCAATTCGATGATGATGATCTAAATCTGTTAAATTTGT 407  
DB 3093 TTTTCTTTAGATAATAGATAATTAATGTTTGTATAGGAAGATAATGATGATGATTTT 3152  
QY 408 GAACA 412  
DB 3153 TAAAA 3157  
RESULT 10  
ID ADB54179  
XX ADB54179 standard; DNA; 5987 BP.  
XX

AC ADB54179;  
 XX  
 DT 04-DEC-2003 (first entry)  
 DE  
 DE Pretreated genomic DNA region 103.  
 XX  
 KW colon cell proliferative disorder; non methylated CpG dinucleotide;  
 KW cytostatic; cancer; adenoma; carcinoma; cytosine methylation state; ds.  
 XX Unidentified.  
 OS  
 XX WO2003072821-A2.  
 FN  
 XX  
 XX 04-SEP-2003.  
 PD  
 XX  
 XX 27-FEB-2003; 2003WO-EP002035.  
 PF  
 XX 27-FEB-2002; 2002EP-00004551.  
 PR  
 XX (EPIG-) EPIGENOMICS AG.  
 XX  
 XX Adorjan P, Burger M, Maier S, Nimmrich I, Becker B, Lesche R;  
 PI Rujan T, Schmitt A;  
 XX  
 XX WPI; 2003-731620/69.  
 DR  
 XX  
 XX Detecting and differentiating between colon cell proliferative disorders  
 PT associated with a gene or its regulatory regions comprises contacting a  
 PT target nucleic acid in a biological sample obtained from the subject with  
 PT a reagent.  
 XX  
 XX Claim 32; SEQ ID NO 235; 74pp; English.  
 PS  
 XX The invention relates to a novel method for detecting and differentiating  
 CC between colon cell proliferative disorders associated with at least one  
 CC gene or its regulatory regions. The method comprises contacting a target  
 CC nucleic acid in a biological sample obtained from the subject with at  
 CC least one reagent or a series of reagents, where the reagent or series of  
 CC reagents, distinguishes between methylated and non methylated CpG  
 CC dinucleotides within the target nucleic acid. The molecules of the  
 CC invention demonstrate cytosine methylation state or single nucleotide  
 CC for detecting and differentiating between colon cell proliferative  
 CC disorders, including cancers such as colon adenoma and colon carcinoma.  
 CC The PNA (peptide nucleic acid)-oligomers are useful as probes for  
 CC determining cytosine methylation state or single nucleotide  
 CC polymorphisms. The current sequence is that of the pretreated genomic DNA  
 CC region of the invention. This sequence is not shown within the  
 CC specification but is taken from Wipoweb.  
 XX  
 XX Sequence 5987 BP; 1754 A; 96 C; 1096 G; 3041 T; 0 U; 0 Other;  
 SQ

Query Match 11.7%; Score 49; DB 9; Length 5987;  
 Best Local Similarity 47.5%; Pred. No. 0.026;  
 Matches 145; Conservative 0; Mismatches 160; Indels 0; Gaps 0;

QY 108 TTCTATCTCGCCCAATATCTGTTCTTCGCGCAATGATCTTCGTAAGATTTTA 167  
 DB 2853 TTTTITTTTATGTTAGATTTGTTATTTTAAAAATTTTGTATTTTAAAGTTTA 2912

QY 168 GGTTAGAGATGAATCGTATAGTATTCATCACCAGATAGTTCTTGTCTAGAACT 227  
 DB 2913 GATTTTITGAAATTTTITTTTGTATTTTAAATATGATGATTTTATTTTATTT 2972

QY 228 CTGAATTCCTGATGATTTTCAATGATGTAATAAGATGTTCTTATTCGCGCATTTGTA 287  
 DB 2973 GTTATATAAGTATGATGAATTTTATTTATGATGATGATGATGATTTTATTTTGA 3032

QY 288 TTAGGTTTGTATTTCTTCTGATATGCGATTGCAATTAGGATTTTCTTGTGTTGTGT 347  
 DB 3033 AAATAAATTAATAATAATTTTATTTTATGTTTGTATTTTATTTTGTATTTAAAT 3092

QY 348 TGATCTTTACGATACATTCCTGCAATGAATACGATACGATCTAAATCTTGTATTTGTT 407

DB 3093 TTTTITTTAGATATAGATAATTAATTAATGTTTGTATAGGAGATAATGATGATTAATT 3152

QY 408 GAACA 412  
 DB 3153 TAAAA 3157

RESULT 11  
 ABK39964  
 ID ABK39964 standard; DNA; 15479 BP.  
 XX  
 XX ABK39964;  
 AC  
 XX 21-MAY-2002 (first entry)  
 DT  
 XX Human chemically pretreated gene sequence #23 strand 1.  
 DE  
 XX Human; ds; bisulphite treatment; CpG; DNA methylation; cancer; tumour;  
 KW cytosine; ALDH6; CYP11A; CYP11B1; CYP3A3; DPYD; EPHX2; OCLN; TXNRD1;  
 KW UGT8; MRP; pharmacogenomics; SNP; single nucleotide polymorphism.  
 XX  
 XX Homo sapiens.  
 OS  
 XX WO200202806-A2.  
 FN  
 XX 10-JAN-2002.  
 PD  
 XX 29-JUN-2001; 2001WO-EP007470.  
 PF  
 XX 30-JUN-2000; 2000DE-01032529.  
 PR 01-SEP-2000; 2000DE-01043826.  
 XX  
 XX (EPIG-) EPIGENOMICS AG.  
 PA  
 XX Olek A, Piepenbrock C, Berlin K;  
 PI  
 XX WPI; 2002-154757/20.  
 DR  
 XX New nucleic acid, oligonucleotides and peptide nucleic acid-oligomers,  
 PT useful for detecting cytosine methylation state of genes associated with  
 PT pharmacogenomics and for therapy of diseases e.g. cancer.  
 XX  
 XX Claim 1; SEQ ID NO 45; 24pp; English.  
 PS  
 XX The invention relates to a nucleic acid comprising a sequence at least 18  
 CC bases in length of a segment of the chemically pretreated DNA of genes  
 CC associated with pharmacogenomics according to one of the sequences of the  
 CC genes ALDH6 (NM 000693), CYP11A (NM 000781), CYP11B1 (NM 000497), CYP3A3  
 CC (NM 000776 and NM 017480), DPYD (NM 000110), EPHX2 (NM 001979), OCLN  
 CC (NM 002538), TXNRD1 (NM 003330), UGT8 (NM 003360), MRP (NM 004996,  
 CC NM 019900, NM 019901, NM 019902, NM 019862, NM 019898, NM 019899) and  
 CC their complementary sequences, or a sequence (S1) chosen from 87  
 CC sequences and their complements. The chemical pretreatment is bisulphite  
 CC treatment to convert cytosines (but not methyl-cytosines) into uracil.  
 CC Also included are an oligomer (II) in particular an oligonucleotide or a  
 CC peptide nucleic acid (PNA)-oligomer, comprising in each case at least one  
 CC base sequence having a length of 9 nucleotides which hybridises to or is  
 CC identical to a chemically pretreated DNA of genes associated with  
 CC pharmacogenomics and their complements, arranged in an array for  
 CC analysing diseases associated with the methylation state (CpG) and/or  
 CC detecting SNPs (single nucleotide polymorphisms) of the 87 sequences. The  
 CC oligomers may also be used as PCR primers. The set of 87 nucleic acids  
 CC and their complements is useful for diagnosis and therapy of solid  
 CC tumours and cancer. The present sequence represents one the 87 DNA  
 CC sequences or its complement. Note: The sequence data for this patent did  
 CC not form part of the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
 XX Sequence 15479 BP; 4303 A; 122 C; 2850 G; 8204 T; 0 U; 0 Other;  
 SQ

Query Match 11.7%; Score 48.8; DB 6; Length 15479;  
 Best Local Similarity 51.9%; Pred. No. 0.034;

Matches 110; Conservative 0; Mismatches 102; Indels 0; Gaps 0;  
Qy 204 AGATAGTTCTTTGTCAGAAATCTCTGAAATCTCGATAGTTTTCACATGTCGTAATAGA 263  
Db 11257 AGGAGTTTCTTTGTTTAAATATTTTGTGATTTTATATTTTGTATATGTCGT 11316  
Qy 264 TTGTTCTTATTCGCGGATCTTGTATAGGTTTGTGATTTTCTTGATATGCGATTCGAT 323  
Db 11317 TTTTATTTATTAAGATGTTTGTGATTTTATTTATATATGTTTATTTATTTATGTCGTT 11376  
Qy 324 TAGGATTTTCTTTGTTTGTGTTGATCTTACATACATCTCTGCAATGAATACGAT 383  
Db 11377 GGGGTTGTTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTT 11436  
Qy 384 GGATCTAAATCTTCTTAAATTTGTTGACAGAT 415  
Db 11437 TGGTTAGATATTATTTATTTTAAAAAGAT 11468

RESULT 12  
AAS45323  
ID AAS45323 standard; DNA; 17848 BP.  
AC AAS45323;  
XX  
DT 18-DEC-2001 (first entry)  
DE Chemically pretreated complementary DNA associated with cell cycle #14.  
KW Cell cycle; human; CpG dinucleotide; cytosine methylation; HIV; aging;  
KW human immunodeficiency virus; neurodegenerative disorder; solid tumour;  
KW graft-versus-host disease; glomerular disease; Lewy body disease; cancer;  
KW arthritis; arteriosclerosis; anti-HIV; neuroprotective; antiarthritis;  
KW immunosuppressive; antitumour; cytostatic; antiarteriosclerotic; ds;  
KW PCR primer.

OS Homo sapiens.  
XX  
XX  
XX WO200168911-A2.  
XX  
XX  
XX 20-SEP-2001.  
XX  
XX 15-MAR-2001; 2001WO-EP002945.  
XX  
XX 15-MAR-2000; 2000DE-01013847.  
XX  
XX 06-APR-2000; 2000DE-01019058.  
XX  
XX 07-APR-2000; 2000DE-01019173.  
XX  
XX 30-JUN-2000; 2000DE-01032529.  
XX  
XX 01-SEP-2000; 2000DE-01043826.  
XX  
XX (EPIG-) EPIGENOMICS AG.  
XX  
XX Olek A, Piepenbrock C, Berlin K;  
XX  
XX WPI; 2001-602751/58.  
XX  
XX Designing primers and probes for analyzing diseases associated with  
XX cytosine methylation state e.g. arthritis, cancer, aging,  
XX arteriosclerosis comprising fragments of chemically modified genes  
XX associated with cell cycle.  
XX  
XX Claim 1; SEQ ID NO 28; 28pp; English.

XX Sequences AAS45296-AAS45520 represent chemically pretreated genomic DNA  
XX molecules associated with the cell cycle and specific PCR primers of the  
XX invention. The sequences are useful for detecting the methylation state  
XX of all CpG dinucleotides in a sequence and therefore for analysing  
XX associated diseases. By analysing cytosine methylations in the pretreated  
XX DNA, genetic and/or epigenetic parameters for the diagnosis and therapy  
XX of existing diseases or the predisposition to specific diseases can be  
XX ascertained. The parameters may be compared to another set of genetic  
XX and/or epigenetic parameters, the differences serving as basis for  
XX diagnosis and/or prognosis events which are disadvantageous to patients.

CC The sequences of the invention are useful for the diagnosis and therapy  
CC of HIV infection, neurodegenerative disorders, graft-versus-host disease,  
CC aging, glomerular disease, Lewy body disease, arthritis,  
CC arteriosclerosis, solid tumours and cancers  
XX  
SQ Sequence 17848 BP; 5055 A; 211 C; 3533 G; 9043 T; 0 U; 6 Other;  
Query Match 11.6%; Score 48.4; DB 4; Length 17848;  
Best Local Similarity 46.9%; Pred. No. 0.044;  
Matches 151; Conservative 0; Mismatches 171; Indels 0; Gaps 0;  
Qy 91 AAATPAGCGAGATGATGTTCTATCTTGGCCATATCTTGTTCCTCGCCGATGATCT 150  
Db 1226 AAATATTTTAAAGTTAATATATTTTTTAAATGAATTAATTTTTTTTTTAAATGAG 1285  
Qy 151 TCCGTATTAAGATTTTACGTTAGATGATGATCGTATAGTTCATCACCAGATAGT 210  
Db 1286 TTTTCTTTTGTATTTTACGTTTACGTTTACGTTTACGTTTACGTTTACGTTTACGTT 1345  
Qy 211 TTTCTTTGTCTAGATCTCTGAAATCTCGATGTTTCCATGTTTCCATGTTTCCATGTTTCT 270  
Db 1346 TTTTCTTTTGTAAATTTTGTGTTTATTTAAATTTTTTTTGTATTTTGTATTTTATTTT 1405  
Qy 271 TATTGCGCGATTTGTTGATGAGGTTTTCATTTTCTTGTATTTGCGATTTGCAATTAGGAT 330  
Db 1406 TATTGTTTCTTTTCTTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTT 1465  
Qy 331 TTTCTTTGTTTGTGTTGATCTTACGATACATCTCTGCAATTTGAAATGATGATGATCTA 390  
Db 1466 TGTCTTTTGTGTTGTTTCTTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTT 1525  
Qy 391 AATCTTGTAAATTTGTTGAAACA 412  
Db 1526 ATATGAATAATATTTGTTTATA 1547

RESULT 13  
ABK39976  
ID ABK39976 standard; DNA; 17848 BP.  
XX  
XX AC ABK39976;  
XX  
XX 21-MAY-2002 (first entry)  
XX  
XX Human chemically pretreated gene sequence #29 strand 2.  
XX  
XX Human; ds; bisulphite treatment; CpG; DNA methylation; cancer; tumour;  
XX cytosine; ALDH6; CYP11A; CYP11B; CYP3A3; DPID; EPHX2; OCLN; TXNRD1;  
XX UGT8; MRP; pharmacogenomics; SNP; single nucleotide polymorphism.  
XX  
XX Homo sapiens.  
XX  
XX WO200202806-A2.  
XX  
XX 10-JAN-2002.  
XX  
XX 29-JUN-2001; 2001WO-EP007470.  
XX  
XX 30-JUN-2000; 2000DE-01032529.  
XX  
XX 01-SEP-2000; 2000DE-01043826.  
XX  
XX (EPIG-) EPIGENOMICS AG.  
XX  
XX Olek A, Piepenbrock C, Berlin K;  
XX  
XX WPI; 2002-154757/20.  
XX  
XX New nucleic acid, oligonucleotides and peptide nucleic acid-oligomers,  
XX useful for detecting cytosine methylation state of genes associated with  
XX pharmacogenomics and for therapy of diseases e.g. cancer.  
XX  
XX Claim 1; SEQ ID NO 58; 24pp; English.

CC The invention relates to a nucleic acid comprising a sequence at least 18  
 CC bases in length of a segment of the chemically pretreated DNA of genes  
 CC associated with pharmacogenomics according to one of the sequences of the  
 CC genes ALDH6 (NM 000693), CYP11A (NM 000781), CYP11B (NM 000497), CYP3A3  
 CC (NM 000776 and NM 017460), DPYD (NM 000110), EPHX2 (NM 001979), OCLN  
 CC (NM 002538), TXNRD1 (NM 003330), UGT8 (NM 003360), MRP (NM 004996),  
 CC NM 019900, NM 019901, NM 019902, NM 019862, NM 019898, NM 019899, and  
 CC their complementary sequences, or a sequence (S<sub>i</sub>) chosen from 87  
 CC sequences and their complements. The chemical pretreatment is bisulphite  
 CC treatment to convert cytosines (but not methyl-cytosines) into uracils.  
 CC Also included are an oligomer (II) in particular an oligonucleotide or a  
 CC peptide nucleic acid (PNA)-oligomer, comprising in each case at least one  
 CC base sequence having a length of 9 nucleotides which hybridises to or is  
 CC identical to a chemically pretreated DNA of genes associated with  
 CC pharmacogenomics and their complements, arranged in an array for  
 CC analysing diseases associated with the methylation state (CpG) and/or  
 CC detecting SNPs (single nucleotide polymorphisms) of the 87 sequences. The  
 CC oligomers may also be used as PCR primers. The set of 87 nucleic acids  
 CC and their complements is useful for diagnosis and therapy of solid  
 CC tumours and cancer. The present sequence represents one the 87 DNA  
 CC sequences or its complement. Note: The sequence data for this patent did  
 CC not form part of the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 17848 BP; 5055 A; 211 C; 3533 G; 9043 T; 0 U; 6 Other;

Query Match 11.6%; Score 48.4; DB 6; Length 17848;  
 Best Local Similarity 46.9%; Pred. No. 0.044;  
 Matches 151; Conservative 0; Mismatches 171; Indels 0; Gaps 0;

QY 91 AAATTAGGCGAGATTAGTTCTTATCTTCTGGCCATATCTTCTTCTGCGCGAATGATCT 150  
 DB 1226 AATATTTTAAAGTTAATATATTTTAAATGAATTAATTTTATTAATTCAG 1285

QY 151 TCGTATAAGATTCTGATAGATGAATCGTATAGATTTTATGATTTTATTAATTCAGT 210  
 DB 1286 TTTTCTTTGTTAAATTTAGTGGTTAATTAATTTTCTGATTTGTTTATTTTATA 1405

QY 271 TATTCGGCGATTGTTAGTGGTTTGTATTTCTTGTATTTGCGAATTCAGATTCAGGAT 330  
 DB 1406 TATTTGTTTCTTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTT 1465

QY 331 TTTCTTTGGTTTGTGTTGATCTTACGATACATCTCGCAATTCAGTATGATCTA 390  
 DB 1466 TGTCTTTTTCATGTTGTTTGTGTTTCTTTTATTTTATTTATTTATTTGTTAGAT 1525

QY 391 AATCTGTTAAATTTGTTGACA 412  
 DB 1526 ATATGAATAAATTTGTTTATA 1547

RESULT 14  
 ABK28164  
 ID ABK28164 standard; DNA; 17848 BP.

XX ABK28164;

XX 23-APR-2002 (first entry)

XX DNA transcription associated complementary genomic DNA #19.

XX DNA transcription associated gene; peptide nucleic acid; PNA-oligomer;  
 KW PNA; cytosine methylation state; SNP; retroviral infection; gene; ds;  
 KW single nucleotide polymorphism; adenosine deaminase deficiency; cancer;  
 KW viral infection; Sezary syndrome; haematological disorder; tuberculosis;  
 KW immunological disorder; Werner syndrome; developmental disorder;  
 KW psoriasis; Rieger's syndrome; neurological disorder; erythropoiesis;  
 KW neurodegenerative disorder; Waardenburg syndrome; Niemann-Pick disease;

KW myelodysplastic syndrome; myocardial infarction; hypertension; arthritis;  
 KW angiogenesis; congenital heart disease; HDR syndrome; gene therapy;  
 KW polyglutamine disorder; solid tumour.

XX Unidentified.

XX W0200192565-A2.

XX 06-DEC-2001.

XX 06-APR-2001; 2001WO-EP003973.

XX 06-APR-2000; 2000DE-01019058.

XX 07-APR-2000; 2000DE-01019173.

XX 30-JUN-2000; 2000DE-01032529.

XX 01-SEP-2000; 2000DE-01043826.

XX (EPIG-) EPICENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K;

XX WPI; 2002-090046/12.

XX New nucleic acids or oligomers, useful for diagnosing or treating  
 XX diseases associated with DNA transcription, e.g. immunological disorders,  
 XX Werner syndrome, psoriasis, myocardial infarction, solid tumors or  
 XX cancer.

XX Claim 1; SEQ ID NO 38; 32pp; English.

XX The invention relates to a nucleic acid, which comprises a segment of the  
 XX chemically pretreated DNA of genes associated with DNA transcription from  
 XX one of 346 sequences, and an oligomer, in particular an oligonucleotide  
 XX or peptide nucleic acid (PNA)-oligomer that hybridises to or is identical  
 XX to the chemically pretreated DNA of genes associated with DNA  
 XX transcription. The set of oligomer probes are useful for detecting the  
 XX cytosine methylation state and/or single nucleotide polymorphisms (SNPs)  
 XX in a chemically pretreated genomic DNA. The nucleic acids are useful for  
 XX diagnosing or treating diseases associated with DNA transcription  
 XX (particularly with the methylation status), e.g. adenosine deaminase  
 XX deficiency, viral infection, retroviral infection, Sezary syndrome,  
 XX haematological disorders, immunological disorders, Werner's syndrome,  
 XX tuberculosis, developmental disorders, psoriasis, Rieger's syndrome,  
 XX neurological disorders, neurodegenerative disorders, Waardenburg  
 XX syndrome, Niemann-Pick disease, myelodysplastic syndrome, myocardial  
 XX infarction, hypertension, angiogenesis, erythropoiesis, congenital heart  
 XX disease, HDR syndrome, arthritis, polyglutamine disorders, solid tumours  
 XX or cancer. Sequences ABK28127-ABK28472 represent DNA transcription  
 XX associated genomic DNA molecules of the invention. Note: The sequence  
 XX data for this patent did not form part of the printed specification but  
 XX was obtained in electronic format directly from the European Patent  
 XX Office

XX Sequence 17848 BP; 5055 A; 211 C; 3533 G; 9043 T; 0 U; 6 Other;

Query Match 11.6%; Score 48.4; DB 6; Length 17848;  
 Best Local Similarity 46.9%; Pred. No. 0.044;  
 Matches 151; Conservative 0; Mismatches 171; Indels 0; Gaps 0;

QY 91 AAATTAGGCGAGATTAGTTCTTATCTTCTGGCCATATCTTGTCTTCTGCGCGAATGATCT 150

DB 1226 AATATTTTAAAGTTAATATATTTTAAATGAATTAATTTTATTTTATTAATTCAG 1285

QY 151 TCGTATAAGATTTTTAGGTTAGATGAATCGTATAGATTTTATGATTTTATTAATTCAGT 210

DB 1286 TTTTCTTTGTTAAATTTAGTGGTTAATTAATTTTCTGATTTTATTTATTTATTCAGAT 1345

QY 211 TTTCTTTGTTAGAAATCTGAAATTTCTGATAGTTTTCACATGTCGTAATAGATTTTCT 270

DB 1346 TTTTCTTTGTTAAATTTTAGTGGTTAATTAATTTTCTGATTTTATTTTATTTTATA 1405

QY 271 TATTCGGCGATTGTTGATTAGGTTTCTGATTTTCTGATTTATGCGAATTCAGGAT 330

```

SQ      Sequence 2501 BP; 532 A; 124 C; 707 G; 1138 T; 0 U; 0 Other;
      Query Match      11.5%; Score 48; DB 7; Length 2501;
      Best Local Similarity 47.1%; Pred. No. 0.04;
      Matches 180; Conservative 200; Indels 2; Gaps 1;

      25  TTGATTTTCCTGAAATATTTTTTCGGTGATCGTGAAACTACTGGAATCGCTCGATAGGT 84
      QY

      1661  TTATATATTTTCGAGATTAGTTTTTCGGTTAAATAGTTTGATTTTAAATGTTCGAAGGA 1720
      Db

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1721	AAAGGAAAGTCTGCTGTTATATTCGGTTTTTTTGTGTTTAGTTTTTTAAATGTGAAA	1780
Db		
145	TGATCTTCCGTATAAAGATTTTAGGTTAGAGATGAATCGTATAGCTAGATTTTCATCACCA	204
Qy		
1781	TGGAGTTCGTAGTGTTTTTTATTCGAGGTATGAAAGTTTTTATAGTTTTTAGTTTAT	1840
Db		
205	GATAGTTCTTCTCTAGAAATCTCGAAATCTCGATAGTTTTTCAATGCTGTAATAGAT	264
Qy		
1841	AATAGTTGTGTTAGTTATTATTATGATGATGATTACGATATGTTATTTTTTGGGGAAA	1900
Db		
265	TGTCCTATTCCGCGAATGTTGATTAGGGTTTGATTTTCTTGATTATCGGATTCGAATT	324
Qy		
1901	TAAATATTTTTTCTTTTATGATAGTATTGGTTTTTTTTTTTTTTT--TCGTATTATTAAT	1958
Db		

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047 225 1959 Db QY Db
      1959  TTGGTTTATTATTTTGGTTTAGTTTTTTTATAGGTGATTGTTGTAGGGGTGA 2018
      385  GATCTAAATCTGTTTAATTGT 406
      2019  GATATTATTGTTTTTGTGTTTTT 2040

Search completed: June 20, 2004, 01:58:56
Job time : 312.833 secs

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C	1	379.8	90.9	436	28	BH633573	BH633573	SALK 0427
C	2	353.2	84.5	366	28	BH846820	BH846820	SALK 0105
C	3	194.4	46.5	258	29	AL950572	AL950572	Arabidops
C	4	192.4	46.0	262	29	AL950573	AL950573	Arabidops

Shankar, V. *Unpublished*  
A Sequence-Indexed Library of Insertion Mutations in the  
Arabidopsis Genome  
Unpublished (2001)  
Contact: Joseph R. Ecker  
Salk Institute Genomic Analysis Laboratory (SIGNAL)  
The Salk Institute for Biological Studies  
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
Tel.: 858 453 4100 x1752  
Fax: 858 556 6379  
Email: [ecker@salk.edu](mailto:ecker@salk.edu)

This is single pass sequence recovered from the left border of  
 TDNA. This sequence lies within an annotated exon of At4g40040.  
 Class: TDNA tagged.

## FEATURES

Location/Qualifiers  
 1. 436  
 /organism="Arabidopsis thaliana"  
 /mol\_type="genomic DNA"  
 /strain="Columbia 0"  
 /db\_xref="taxon:3702"  
 /clone="SALK\_042781"  
 /clone\_lib="Arabidopsis thaliana TDNA insertion lines"  
 /note="PCR was performed on Arabidopsis thaliana lines  
 each of which contains one or more TDNA insertion  
 elements. The resultant fragment for each line was  
 directly sequenced to determine the genomic sequence at  
 the site of insertion. Details of the protocols used can  
 be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)"

## ORIGIN

Query Match 90.9%; Score 379.8; DB 28; Length 436;  
 Best Local Similarity 99.5%; Pred. No. 1.6e-67;  
 Matches 381; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 33 CCTCGAATATTTTTCGGTATCGTGAATCTGGAATCGCTCGATAGGTGTCAGAA 92  
 435 CCTGGAATATTTTTCGGTATCGTGAATCTGGAATCGCTCGATAGGTGTCAGAA 376  
 93 ATTAGGCGAGATTAGTTCTATTCTTGGCCGATATCTGTTCTTCCGCAATGATCTTC 152  
 375 ATTAGGCGAGATTAGTTCTATTCTTGGCCGATATCTGTTCTTCCGCAATGATCTTC 316  
 153 CGTATAAGATTAGGTTAGAGATGAATCGTATAGTAGTATCTATCACCAGATGTTT 212  
 315 CGGATAAGATTAGGTTAGAGATGAATCGTATAGTAGTATCTATCACCAGATGTTT 256  
 213 CTTTGTCTAGAACTCTCGAATTCGCAATGTTTCACATGTAATAGATTCTTCTTA 272  
 255 CTTTGTCTAGAACTCTCGAATTCGCAATGTTTCACATGTAATAGATTCTTCTTA 196  
 273 TTCGCGGATCTGATTAGGTTTGTATTTCTTGGATTATCGGATTCGCAATAGGATTT 332  
 195 TTCGCGGATCTGATTAGGTTTGTATTTCTTGGATTATCGGATTCGCAATAGGATTT 136  
 333 TCTTTGGTTTGTGTGTATCTTACGATACATCTCCGCAATGAATGATATGATCTAAA 392  
 135 TCTTTGGTTTGTGTGTATCTTACGATACATCTCCGCAATGAATGATATGATCTAAA 76  
 393 TCTTTGGTTTGTGTGTATCTTACGATACATCTCCGCAATGAATGATATGATCTAAA 415  
 75 TCTTTGGTTTGTGTGTATCTTACGATACATCTCCGCAATGAATGATATGATCTAAA 53

RESULT 2  
 BH46820/c 366 bp DNA linear GSS 13-JUN-2002  
 LOCUS SALK\_010583.54.50.x Arabidopsis thaliana TDNA insertion lines  
 DEFINITION Arabidopsis thaliana genomic clone SALK\_010583.54.50.x, genomic  
 survey sequence.

ACCESSION BH46820

VERSION BH46820.1

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM

Arabidopsis thaliana  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.  
 1 (bases 1 to 366)

REFERENCE

AUTHORS

Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R.,  
 Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L.,  
 Shinn, P., Zimmerman, J. and Ecker, J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished (2001)

TITLE

JOURNAL

## COMMENT

Contact: Joseph R. Ecker  
 Salk Institute Genomic Analysis Laboratory (SIGNAL)  
 The Salk Institute for Biological Studies  
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA  
 Tel: 858 453 4100 x1752  
 Fax: 858 558 6379  
 Email: ecker@salk.edu  
 This is single pass sequence recovered from the left border of  
 TDNA. This sequence lies within 300 bases of the 5' end of  
 At4g40040.  
 Class: TDNA tagged.

## FEATURES

Location/Qualifiers  
 1. 366  
 /organism="Arabidopsis thaliana"  
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 /db\_xref="taxon:3702"  
 /clone="SALK\_010583.54.50.x"  
 /clone\_lib="Arabidopsis thaliana TDNA insertion lines"  
 /note="PCR was performed on Arabidopsis thaliana lines  
 each of which contains one or more TDNA insertion  
 elements. The resultant fragment for each line was  
 directly sequenced to determine the genomic sequence at  
 the site of insertion. Details of the protocols used can  
 be found at [http://signal.salk.edu/tdna\\_protocols.html](http://signal.salk.edu/tdna_protocols.html)"

## ORIGIN

Query Match 84.5%; Score 353.2; DB 28; Length 366;  
 Best Local Similarity 97.8%; Pred. No. 4.1e-62;  
 Matches 358; Conservative 0; Mismatches 8; Indels 0; Gaps 0;  
 30 TTCTCTGGAATATTTTTCGGTATCGTGAATCTGGAATCTCGATAGGTGTCAC 89  
 366 TTCTCTGGAATATTTTTCGGTATCGTGAATCTGGAATCTCGATAGGTGTCAC 307  
 90 GAAATTTAGCGAGATTAGTTCTATTCTTGGCAATATCTTCTTCCGCAATGATC 149  
 306 GAAATTTAGCGAGATTAGTTCTATTCTTGGCAATATCTTCTTCCGCAATGATC 247  
 150 TTCCGATTAAGATTAGTTAGTAGAGTAATCGTATAGTAGTATCTATCACCAGATG 209  
 246 TTCCGATTAAGATTAGTTAGTAGAGTAATCGTATAGTAGTATCTATCACCAGATG 187  
 210 TTCTTTTGTCTAGATCTCTGAAATCTCGATAGTTTTCACATGTTAAATAGATTGTC 269  
 186 TTCTTTTGTCTAGATCTCTGAAATCTCGATAGTTTTCACATGTTAAATAGATTGTC 127  
 270 TTATTCGCGGATTTGTTGATTAGGTTTTCGATTTTCTGATTATGCGATTGCAATTAGGGA 329  
 126 TTATTCGCGGATTTGTTGATTAGGTTTTCGATTTTCTGATTATGCGATTGCAATTAGGGA 67  
 330 TTCTCTTGGTTTGTGTTGATCTTACGATACATCTCCGCAATGGAATGATGATCT 389  
 66 TTCTCTTGGTTTGTGTTGATCTTACGATACATCTCCGCAATGGAATGATGATG 7  
 390 AAATCT 395  
 6 AAATCT 1

## RESULT 3

AL950572/c

LOCUS

AL950572 258 bp DNA linear GSS 24-OCT-2002  
 DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-329E10-016035,  
 genomic survey sequence.

ACCESSION AL950572

VERSION AL950572.1

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM

Arabidopsis thaliana  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

[illegible]





Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 1201)  
Li, W.-S., Gruber, C., Jessee, J., and Polayes, D.  
Full-length cDNA libraries and normalization  
Unpublished (2001)  
On Feb 13, 2001 this sequence version replaced gi:12788300.  
Contact: Genoscope  
Genoscope - Centre National de Sequencage  
BP 191 91006 EVRY cedex - France  
Email: segref@genoscope.cns.fr, Web: www.genoscope.cns.fr  
Library was constructed by life technologies, a division of  
Invitrogen. This sequence belongs to sequence cluster 3928.f. For  
more information about this cluster, see  
http://www.genoscope.cns.fr/  
cgi-bin/cluster.cgi?seq=CS0DC008AF01NP1&cluster=3928.f. Contact :  
Feng liang Email : fliang@lifetech.com URL :  
http://fulllength.invitrogen.com/ Invitrogen Corporation 1600  
Faraday Avenue Genoscope sequence ID : CS0DC008AF01NP1.  
Location/Qualifiers  
1..1201

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT

FEATURES  
source

/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="CS0DC008YK01"  
/issue\_type="NEUROBLASTOMA COT 25-NORMALIZED"  
/clone\_lib="Homo sapiens NEUROBLASTOMA COT 25-NORMALIZED"  
/note="1st strand cDNA was primed with a NotI-oligo(dT)  
primer. Five prime end enriched, double-strand cDNA was  
digested with Not I and cloned into the Not I and EcoR V  
sites of the pCMVSPORT 6 vector. Library was normalized."

ORIGIN

Query Match 12.8%; Score 53.4; DB 9; Length 1201;  
Best Local Similarity 42.9%; Pred. No. 0.3;  
Matches 129; Conservative 23; Mismatches 149; Indels 0; Gaps 0;  
QY 104 TTAGTTCTTATCTGGCATTATCTCTTCTTCGCGGAATGATCTTCGGTATAAAGAT 163  
DB 203 TTTTCTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 262  
QY 164 TTAGGTTAGAGATGAATCGTATAGTATGATTCATACACAGATGATTTCTTTGTTT 223  
DB 263 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 322  
QY 224 ATCTCTGAATTCCTGATAGTTTTCACATGCTAAATAGATTTCTTTATTCGCGGATTG 283  
DB 323 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 382  
QY 284 TTGATTAGGTTTCTGATTTCTTGAATGCGAATGCAATGAGGATTTCTTTGTTT 343  
DB 383 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 442  
QY 344 GTGTTGATCTACGATACATTCCTCGAATGCAATGCAATGCAATGCAATGCAATGCAAT 403  
DB 443 WTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 502  
QY 404 T 404  
DB 503 W 503

RESULT 10  
CNS005F3/c  
LOCUS  
DEFINITION  
Drosophila melanogaster genome survey sequence TET3 end of BAC #  
BACR1P07 of RPCL-98 library from Drosophila melanogaster (fruit  
fly), genomic survey sequence.  
ACCESSION  
AL059925.1 GI:4943047  
VERSION  
GSS.  
KEYWORDS  
Drosophila melanogaster (fruit fly)  
SOURCE  
Drosophila melanogaster  
ORGANISM

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidea; Drosophilidae; Drosophila.  
1 (bases 1 to 960)

REFERENCE  
AUTHORS  
TITLE  
JOURNAL

COMMENT

Direct Submission  
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :  
BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr)  
- Web : www.genoscope.cns.fr  
Determination of this BAC-end sequence was carried out as part of a  
collaboration with the Berkeley Drosophila Genome Project (BDGP).  
The BDGP is constructing a physical map of the Drosophila  
melanogaster genome using these BACs. For further information  
please see http://www.fruitfly.org The BDGP Drosophila  
melanogaster BAC library was prepared by Kazutoyo Osoegawa and  
Aron Mammoser in Pieter de Jong's laboratory in the Department of  
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,  
NY. The library is named RPCL-98 and was constructed by partial  
EcoRI digestion of Drosophila DNA provided by the BDGP from the  
isogenic strain Y2; cn bw sp, the same strain used for the BDGP's  
P1 and EST libraries. A more detailed description of the library  
and how to order individual BAC clones, the entire library, or  
filters for hybridization from the BACPAC Resource Center can be  
found at http://bacpac.med.buffalo.edu/drosophila\_bac.htm.

FEATURES  
source

Location/Qualifiers  
1..960  
/organism="Drosophila melanogaster"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:7227"  
/clone="BACR1P07"  
/clone\_lib="RPCL-98"  
/note="end : TET3"

ORIGIN

Query Match 12.6%; Score 52.6; DB 29; Length 960;  
Best Local Similarity 40.2%; Pred. No. 0.45;  
Matches 158; Conservative 30; Mismatches 205; Indels 0; Gaps 0;  
QY 15 TCATCTCTTTGATTTTCTCGGAATATTTTTCGGTATCGTGAACACTCTGGAATCG 74  
DB 572 TWGARTKTKTGTGTTTCTGTTTCTGTTTCTGTTTCTGTTTCTGTTTCTGTTTCTGTT 513  
QY 75 CTCGATAGTGGTACGAATAGGCGAGATAGTTCTTATTCCTGGCCATTAATCTGTT 134  
DB 512 TGWRGTGTGTGRADAAATTTKTGAWDGGTGTGTTGTTTCTGTTGTTGTTGTTGTTGTT 453  
QY 135 CTTCCGCGAATGATCTCCGTATAAGATTTAGGTTAGATGATGATCGTATAGCTAGAT 194  
DB 452 KGRKTKGTGRDGTGTTTCTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTT 393  
QY 195 TTCATCACCAGATAGTTTCTTGTCTAGAAATCTCGAAATCTCGATAGTTTTCACATG 254  
DB 392 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 333  
QY 255 GTAATAGATGTTCTTATTCGCGAATGTTGATAGGTTTGTAGTTTGTAGTTTGTAGT 314  
DB 332 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 273  
QY 315 GATTGCAATAGGATTTTCTTGTGTTTGTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTT 374  
DB 272 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 213  
QY 375 AATACGTATGATCTAAATCTTGTAAATTTGTT 407  
DB 212 TTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTTTCTTT 180

RESULT 11  
CNS0081L/c  
LOCUS  
DEFINITION  
Drosophila melanogaster genome survey sequence TET3 end of BAC #  
BACR16N08 of RPCL-98 library from Drosophila melanogaster (fruit  
fly), genomic survey sequence.

ACCESSION AL051208  
 VERSION AL051208.1 GI:4933161  
 KEYWORDS GSS.  
 SOURCE Drosophila melanogaster (fruit fly)  
 ORGANISM Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

# REFERENCE

1 (bases 1 to 832)  
 Genoscope.  
 Direct Submission  
 Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr)

# COMMENT

- Web : www.genoscope.cns.fr  
 Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osogawa and Aaron Mammeter at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain Y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at [http://bacpac.med.buffalo.edu/drosophila\\_bac.htm](http://bacpac.med.buffalo.edu/drosophila_bac.htm).

# FEATURES

Location/Qualifiers  
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 /db\_xref="taxon:7227"  
 /clone="BACR16N08"  
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 /note="end : TET3"

# ORIGIN

Query Match 12.5%; Score 52.4; DB 29; Length 832;  
 Best Local Similarity 39.4%; Pred. No. 0.51;  
 Matches 86; Conservative 38; Mismatches 94; Indels 0; Gaps 0;  
 QY 190 TAGATTTCATCCAGATAGTCTTTCTGTCAGATCTCTGAATCTCGATAGTTTCA 249  
 DB 278 TACTTWTAT 219  
 QY 250 CATGTGTAATAGATGTTCTTATTCGGGATGTTGATAGGTTTTCGATTTCTGAT 309  
 DB 218 TTATTCAT 159  
 QY 310 TAGCGAATTCGAATGAGGATTTCTTTGGTTTGTGTTGATCTTACGATATCTCGC 369  
 DB 158 TTTTAT 99  
 QY 370 AATTGAATGATGATGATCTAAATCTGTAATTTGTT 407  
 DB 98 TTKTAWCYKGTTCATCTAAKATWTWKATYKXKW 61

# RESULT 12

EX416727  
 LOCUS BX416727 Homo sapiens NEUROBLASTOMA 712 bp mRNA linear EST 15-MAY-2003  
 DEFINITION CS0DA011Y114 5-PRIME, mRNA sequence.

# ACCESSION

EX416727  
 VERSION BX416727.1 GI:30765629  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens

# REFERENCE

1 (bases 1 to 712)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

# AUTHORS

Li, W. B., Gruber, C., Jesse, J. and Polayes, D.  
 Full-length cDNA libraries and normalization  
 Unpublished (2001)  
 JOURNAL

# COMMENT

Contact: Genoscope  
 Genoscope - Centre National de Sequencage  
 BP 191 91006 EVRY cedex - France  
 Email: segref@genoscope.cns.fr, Web : www.genoscope.cns.fr  
 Library was constructed by Life Technologies, a division of  
 Invitrogen. Contact : Feng Liang Email : fliang@lifetech.com URL :  
<http://fulllength.invitrogen.com/> Invitrogen Corporation 1600  
 Faraday Avenue Genoscope sequence ID : CS0DA011BE07QPI.

# FEATURES

Location/Qualifiers  
 1..712  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="CS0DA011Y114"  
 /tissue\_type="NEUROBLASTOMA"

/notes="Vector: pcwSPORT 6; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-strand cDNA was digested with Not I and cloned into the Not I and EcoRV sites of the pcwSPORT 6 vector. Library was not normalized."

# ORIGIN

Query Match 12.4%; Score 51.8; DB 13; Length 712;  
 Best Local Similarity 10.8%; Pred. No. 0.69;  
 Matches 33; Conservative 152; Mismatches 120; Indels 0; Gaps 0;  
 QY 104 TTAGTTCTATTCTTGGCAATATCTGTTTCTTCCGCAATGATCTTCGATATAAGAT 163  
 DB 377 TTTTCTTTTWT 436  
 QY 164 TTTAGTTAGATGAATCGTATAGTAGTATCATCCAGATAGTTCTTTGTCTAGA 223  
 DB 437 KKKTKTKKKTK 496  
 QY 224 ATCTCGAATCTCGATAGTTTTCATGTCGTAATAGATGTTCTTATTCGGCGATTG 283  
 DB 497 KKK 556  
 QY 284 TTGATTAGGTTTGTGATTTCTTGTATGTCGATTGCAATTAGGATTTCTTTGGTTT 343  
 DB 557 KKK 616  
 QY 344 GTGTGATCTTACGATACATTCCTCGAATGAAATAGATGGAATCTTAATCTTTAAT 403  
 DB 617 KKK 676  
 QY 404 TGTTG 408  
 DB 677 KKKKK 681

# RESULT 13

CNS0026Z  
 LOCUS CNS0026Z/c  
 DEFINITION Drosophila melanogaster, genome survey sequence 17 end of BAC BACN01A10 of DrosBAC library from Drosophila melanogaster (fruit fly), genomic survey sequence.

# ACCESSION

AL097301  
 VERSION AL097301.1 GI:5608912  
 KEYWORDS GSS.  
 SOURCE Drosophila melanogaster (fruit fly)  
 ORGANISM Drosophila melanogaster

# REFERENCE

1 (bases 1 to 1101)  
 Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.  
 Genoscope.  
 Direct Submission  
 Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :

BP 191 91006 EVRY cedex - FRANCE (E-mail : [segrif@genoscope.cns.fr](mailto:segrif@genoscope.cns.fr))  
- Web : [www.genoscope.cns.fr](http://www.genoscope.cns.fr))  
Determination of this BAC-end sequence was carried out as part of a  
collaboration with the European Drosophila Genome Project (EDGP) -  
<http://www.edgp.ebi.ac.uk> - This Drosophila melanogaster BAC  
library (Dros BAC) was made by Alain Billaud at CEPH (Centre  
d'Etude du Polymorphisme Humain) with funding provided by a WEC  
project grant. The DNA was prepared from embryos by Alain Bucheton  
and Genevieve Payan. It has been constructed in the vector  
pBelOBAC11.

```

FEATURES             Location/Qualifiers
     1..1101
     /organism="Proscophila melanogaster"
     /mol_type="genomic DNA"
     /db_xref="taxon:7227"
     /clone="BACN01A10"
     /clone_lib="DrosBAC"
     /plasmid="pBelBAC11"
     /note="end : T7"

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[illegible]

RESULT 14	CNS0198G	894 bp	DNA	linear	GSS 26-JUL-1999
LOCUS	CNS0198G/c				
DEFINITION	Drosophila melanogaster genome survey sequence T7 end of BAC BAC131816 of DrosBAC library from Drosophila melanogaster (fruit fly), genomic survey sequence.				
ACCESSION	AF109126				
VERSION	AF109126.1	GI:5629430			
KEYWORDS	GSS.				
SOURCE	Drosophila melanogaster (fruit fly)				
ORGANISM	Drosophila melanogaster				
	Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydraeidae; Drosophilidae; Drosophila.				
REFERENCE	1 (bases 1 to 894)				
AUTHORS	Genoscope.				
TITLE	Direct Submission				
JOURNAL	Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : <a href="mailto:secre@genoscope.cns.fr">secre@genoscope.cns.fr</a> )				

- Web : [www.genoscope.cns.fr](http://www.genoscope.cns.fr)  
Determination of this BAC-end sequence was carried out as part of a collaboration with the European Drosophila Genome Project (EDGP) - <http://www.edgp.ebi.ac.uk> - this *Drosophila melanogaster* BAC library (Dros BAC) was made by Alain Billaud at CEPH (Centre d'Etude du Polymorphisme Humain) with funding provided by a MRC project grant. The DNA was prepared from embryos by Alain Bucheton and Genevieve Payan. It has been constructed in the vector pBeloBAC11.

```
FEATURES
source
Location/Qualifiers
1..894
organism="Procephala melanogaster"
mol_type="genomic DNA"
db_xref="taxon:7227"
clone="BACN13E16"
clone_lib="DrosBAC"
plasmid="pBeloBAC11"
notes="end : T7"
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## ORIGIN

Query Match	12.0%;	Score	50.2;	DB	29;	Length	894;
Best Local Similarity	38.1%;	Fred. No.	1.4;				
Matches	140;	Conservative	39;	Mismatches	188;	Indels	0;
						Gaps	0;
Qy	43	TTTTTCGGTGATCGTGAACACTACTGAAATCGGTCCGATAGTGGTACGAAATAGGCGAG	102				
Db	550	TTTTTCTGKTTTTTTTTTAGTCGGKGTGTCACACKTAKYYTKWTATGTTTGTAKAA	491				
Qy	103	ATTAGTTTCPTATCTTGGCCATTATCTTGTTTCCTTCGCCGAATGATCTTCGGTATAAAGA	162				
Db	490	TCATTTTGWGATACGTTCTCTTGATCGWGTATATTTTTGTTTTTKTTTTGTTAKWGT	431				
Qy	163	TTTTAGGTTAGAGATGAATCGTATAGTACATATTTCAACACAGATAGTTCTTTGTCTAG	222				
Db	430	TWTTTTTKTTTAAATCYKWWTTTTYKKTATATTTTTTTTTTKTCWKKTKTTTTYCCAGYTT	371				
Qy	223	AATCTCTGAAATCTCGATAGTTTTACATGCTGTPAAATAGATTCCTTTATCGCGCAAT	282				
Db	370	TGTTTTTGTTTTTTTTTKTTTTTGTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	311				
Qy	283	GTCGATTAGGGTTTTGATTTTCTTGATTATGCGAATTGCAATTAGGGATTTTCTTTGGTTT	342				
Db	310	TTTTTTTTTTGTGTGTGTTTKTTT	251				
Qy	343	TGTCGTGATCTTACGATACATTCCTCGAAATGAATAGTATGAGATCTAAATCTTGTTAAT	402				
Db	250	TBTXYKGTTTTTTKTTGTGWKTTTTYYTTTWWKATWWAKTKAYGTTWTTTKTKTWKTWTTW	191				
Qy	403	TTGTGA	409				
Db	190	TATATKW	184				

RESULT 15  
CNS014RT/C

CNS014RT//C  
LOCUS  
DEFINITION  
CNS014RT  
Drosophila melanogaster genome survey sequence 17 end of SAC  
BACN12L12 of DrosBAC library from Drosophila melanogaster (fruit  
fly), genomic survey sequence.  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :  
BP 191 91006 EVRY cedex - FRANCE (E-mail : sefre@genoscope.cns.fr  
- Web : www.genoscope.cns.fr)





GenCore version 5.1.6  
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: June 20, 2004, 01:39:08 ; Search time 2448.33 Seconds  
(without alignments)  
6025.290 Million cell updates/sec

Title: US-09-000-062-7

Perfect score: 494

Sequence: 1 ctccagggaagaacaggat.....cgcatccgggactctgcg 494

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 27513289 seqs, 14931090276 residues

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:

1: em\_estba.\*

2: em\_esthum.\*

3: em\_estin.\*

4: em\_estmu.\*

5: em\_estov.\*

6: em\_estpl.\*

7: em\_estro.\*

8: em\_hcc.\*

9: gb\_est1.\*

10: gb\_est2.\*

11: gb\_hcc.\*

12: gb\_est3.\*

13: gb\_est4.\*

14: gb\_est5.\*

15: em\_estfun.\*

16: em\_estom.\*

17: em\_gss\_hum.\*

18: em\_gss\_inv.\*

19: em\_gss\_pln.\*

20: em\_gss\_vrt.\*

21: em\_gss\_fun.\*

22: em\_gss\_mam.\*

23: em\_gss\_mus.\*

24: em\_gss\_pro.\*

25: em\_gss\_rod.\*

26: em\_gss\_pig.\*

27: em\_gss\_vxl.\*

28: gb\_gss1.\*

29: gb\_gss2.\*

Result No.	Score	Query Match	Length	DB ID	Description
1	266.4	53.9	300	9	AV551146
2	202.8	41.1	320	14	Z34742 ATTS3502 St
3	81.4	12.4	712	13	BX416727 BX416727
4	54.4	11.0	1101	29	CNS0120N

#### SUMMARIES

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

5	54	10.9	576	29	CNS035N7	AL228940 Tetraodon
C 6	52.4	10.6	1131	14	CD050625	AGENCOURT
C 7	51.8	10.5	683	29	CNS0028F	AL097353 Drosophil
C 8	51.8	10.5	942	29	CNS018GS	AL109318 Drosophil
C 9	51.4	10.4	873	13	BUS34806	BUS34806 AGENCOURT
C 10	51.2	10.4	750	29	CNS011ID	AL100303 Drosophil
C 11	51	10.3	1246	29	CG744146	CG744146 P036-3-E1
C 12	50	10.1	741	29	CNS007YN	AL051401 P043-3-E1
C 13	49.6	10.0	563	29	CNS007UZ	AL051476 Drosophil
C 14	49.6	10.0	737	29	CNS008BU	CG749401 P051-4-D1
C 15	49.4	10.0	1403	29	CG756681	CG756681 P051-4-D1
C 16	49.4	10.0	1388	29	CG756681	CG756681 P051-4-D1
C 17	49.2	10.0	1022	13	BUS29542	BUS29542 AGENCOURT
C 18	49	9.9	1108	29	CNS00710	AL067316 Drosophil
C 19	49	9.9	1201	9	AL514421	AL514421 Tetraodon
C 20	48.8	9.9	483	29	CNS02CV8	AL191645 Tetraodon
C 21	48.8	9.9	848	12	BG809697	BG809697 mgct001xf
C 22	48.8	9.9	1761	28	CC188336	CC188336 CH261-70H
C 23	48.6	9.8	1103	28	BZ551529	BZ551529 pac81-60
C 24	48.6	9.8	1201	13	BX336467	BX336467 BX336467
C 25	48.4	9.8	864	29	CNS0605G	AL07642 T7 end of
C 26	48.4	9.8	1058	14	CD048666	CD048666 AGENCOURT
C 27	48.4	9.8	1300	12	BM468018	BM468018 AGENCOURT
C 28	48.2	9.8	878	29	CNS0028X	AL097371 Drosophil
C 29	48.2	9.8	1059	29	CNS0022B	AL097133 Drosophil
C 30	48.2	9.8	1376	29	CG747831	CG747831 P041-3-B0
C 31	47.8	9.7	922	29	CNS0073W	AL066784 Drosophil
C 32	47.8	9.7	1180	13	BX436369	BX436369 BX436369
C 33	47.8	9.7	1215	12	BM322560	BM322560 AGENCOURT
C 34	47.6	9.6	446	12	BP516575	BP516575 BP516575
C 35	47.6	9.6	914	13	BUS63375	BUS63375 AGENCOURT
C 36	47.6	9.6	967	29	CNS0772W	AL433362 T7 end of
C 37	47.4	9.5	614	29	CNS0152H	AL104915 Drosophil
C 38	47	9.5	867	29	CNS0054A	AL057618 Drosophil
C 39	47	9.5	1053	13	BUS08694	BUS08694 AGENCOURT
C 40	47	9.5	1289	29	CG748984	CG748984 P043-1-C0
C 41	47	9.5	1345	29	CG746697	CG746697 P040-1-B0
C 42	46.8	9.5	767	29	CNS00AQX	AL055924 Drosophil
C 43	46.8	9.5	854	29	CNS012CM	AL101392 Drosophil
C 44	46.8	9.5	966	14	CD388802	CD388802 AGENCOURT
C 45	46.8	9.5	995	14	CD385043	CD385043 AGENCOURT

#### ALIGNMENTS

AV551146 300 bp mRNA linear EST 06-SEP-2000  
AV551146 Arabidopsis thaliana roots Columbia Arabidopsis thaliana  
CDNA clone RZ121b02R 5', mRNA sequence.

AV551146  
AV551146.1 GI:8722559

Arabidopsis thaliana (thale cress)

Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi

1 (bases 1 to 300)

Asamizu, E., Nakamura, Y., Sato, S. and Tabata, S.  
A large scale analysis of cDNA in Arabidopsis thaliana: Generation  
of 12,028 non-redundant expressed sequence tags from normalized and  
size-selected cDNA libraries  
DNA Res. 7 (3), 175-180 (2000)

20363093

10907847

CONTACT: Erika Asamizu

The First Laboratory for Plant Gene Research

Kazusa DNA Research Institute

Yana 1532-3, Kisarazu, Chiba 292-0812, Japan

Email: asamizu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.

Location/Qualifiers

FEATURES

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source
1. 300
/organism="Arabidopsis thaliana"
/mol_type="mRNA"
/db_xref="taxon:3702"
/clone="R2121h02R"
/tissue="roots"
/clone_lib="Arabidopsis thaliana roots Columbia"
/notes="vector: pBluescriptII SK-, Site_1: EcoRI; Site_2: XhoI"

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Matches 267; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

ORIGIN
QY 1 CTCAGGCGAAGACAGATGATGTTGTTCTTAATTAGATCAGGGGTTTAGGCTCTTTCCAT 60
DB 33 CTCAGGCGAAGACAGATGATGTTGTTCTTAATTAGATCAGGGGTTTAGGCTCTTTCCAT 92
QY 61 TACTTTTTTAATGTTTTTCTGTTACTCTCTCCGGATCTGATTTTACGACAATAGAGTTT 120
DB 93 TACTTTTTTAATGTTTTTCTGTTACTCTCTCCGGATCTGATTTTACGACAATAGAGTTT 152
QY 121 CGGGTTTTCCTCCATTCAGTTTGAATAAATGCTCGTCTTTTAAAGTTTTCGTCGATCGA 180
DB 153 CGGGTTTTCCTCCATTCAGTTTGAATAAATGCTCGTCTTTTAAAGTTTTCGTCGATCGA 212
QY 181 TAAACCTGTGAAGATTGAGTCTAGTCGATTTATTGGATGATCCATTTCTTCATCGTTTTT 240
DB 213 TAAACCTGTGAAGATTGAGTCTAGTCGATTTATTGGATGATCCATTTCTTCATCGTTTTT 272
QY 241 TCTTGCTTCGAAGTTCTGTATAACAGA 268
DB 273 TCTTGCTTCGAAGTTCTGTATAACAGA 300

RESULT 2
Z34742
LOCUS
DEFINITION
ATTS3502 Strasbourg-A Arabidopsis thaliana cDNA clone Fall85 5',
mRNA sequence.
ACCESSION
Z34742
VERSION
Z34742.1 GI:507089
KEYWORDS
EST.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 320)
CNRS.
The Arabidopsis thaliana transcribed genome: the GDR cDNA program
Unpublished (1996)
Contact: Philipps G., Gigot C.
Gigot Claude / L512
Laboratoire de Biologie Moleculaire des Plantes - CNRS
12 Rue du General Zimmer, 67084 Strasbourg Cedex, France
Email: ARABANK@MBOC.U-STRASBG.FR.
Location/Qualifiers
1. 320
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/mol_type="mRNA"
/strains="ecotype Columbia"
/db_xref="taxon:3702"
/clone="FA1185"
/clone_lib="Strasbourg-A"
/notes="vector: Lambda ZAPII; tissue-sliced leaves of
A.thaliana ecotype Columbia; clone library=Strasbourg-A;
Cloning vector: Lambda ZAPII; Physiological condition:
leaves strips incubated 2/3/4 days in liquid culture
medium."

FEATURES
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/organism="Homo sapiens"
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/clone_lib="Homo sapiens NEUROBLASTOMA"
/notes="Vector: pCMVSPORT 6; 1st strand cDNA was primed
with a NotI-oligo(dT) primer. Five prime end enriched,
double-strand cDNA was digested with Not I and cloned into
the Not I and EcoRV sites of the pCMVSPORT 6 vector.
Library was not normalized."

ORIGIN
Query Match
Best Local Similarity 12.4%; Score 61.4; DB 13; Length 712;
Matches 53; Conservative 179; Mismatches 165; Indels 0; Gaps 0;

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Query Match
Best Local Similarity 41.1%; Score 202.8; DB 14; Length 320;
Matches 273; Conservative 0; Mismatches 42; Indels 5; Gaps 4;

QY 17 GTATGATTTGTTTCTTAATTAGATCAGGGGTTTAGGCTCTTTCCATTTACTTTTAAAG-TTT 75
DB 1 GTATGATTTGTTTCTTAATTAGATCAGGGGTTTAGGCTCTTTCAATTTACTTTTAAAGTTT 60
QY 76 TTTCTGTTTACTGTCTCC--GCGATCTGATTTTACGACAATAGAGTTTTCGGGTTTGTCCC 133
DB 61 TTTCTGTTTACTGTCTCCGGCGATCTGATTTTACCGCAATAGAGTTTTCGGGTTTGTCCA 120
QY 134 ATTCAGTTTGAATAAATACGT-CGGCTCTTTTAACTTTGCTGATCGATCAAAACCTGTGAA 192
DB 121 TTCCAAGTTTGAATAAATGAGTCCCGCTCTTTTAAAGTTTTCGTCGATCGATCAAAACCTGTGAA 180
QY 193 GATTGAGTCTAGTCTGATTTATTGGATGATCCATTTCTTCATCGTTTTTTTCTTGTTCGAA 252
DB 181 GATTGAGTCTAGTCTGATTTATTGGATGATCCATTTCTTCATCGTTTTTTTCTTGTTCGAA 239
QY 253 GTTCTGTATAACAGATTTGCTGTGCGGATTTCTTACTAGCTAGCCGTGTATCGAGAAC 312
DB 240 GTTCTGTATAACAGATTTGCTGTGCGGATTTCTTACTAGCTAGCCGTGTATCGAGAAC 299
QY 313 TAGGGTTTTTCGAGTCAATTT 332
DB 300 TAGGGTTTTTCGAGTCAATTT 319

RESULT 3
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LOCUS
DEFINITION
BX416727 Homo sapiens NEUROBLASTOMA Homo sapiens cDNA clone
CSODA011Y114 5-PRIME, mRNA sequence.
ACCESSION
BX416727
VERSION
BX416727.1 GI:30765629
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 712)
Li, W.B., Gruber, C., Jessee, J. and Polayes, D.
Full-length cDNA libraries and normalization
Unpublished (2001)
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Web : www.genoscope.cns.fr
Email: seqref@genoscope.cns.fr,
Library was constructed by Life Technologies, a division of
Invitrogen. Contact : Feng Liang Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com/
Invitrogen Corporation 1600
Paradise Avenue Genoscope sequence ID : CSODA011BE07QP1.
Location/Qualifiers
1. 712
/organism="Homo sapiens"
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/db_xref="taxon:9606"
/clone="CSODA011Y114"
/tissue type="NEUROBLASTOMA"
/clone_lib="Homo sapiens NEUROBLASTOMA"
/notes="Vector: pCMVSPORT 6; 1st strand cDNA was primed
with a NotI-oligo(dT) primer. Five prime end enriched,
double-strand cDNA was digested with Not I and cloned into
the Not I and EcoRV sites of the pCMVSPORT 6 vector.
Library was not normalized."

FEATURES
source
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/organism="Homo sapiens"
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/notes="Vector: pCMVSPORT 6; 1st strand cDNA was primed
with a NotI-oligo(dT) primer. Five prime end enriched,
double-strand cDNA was digested with Not I and cloned into
the Not I and EcoRV sites of the pCMVSPORT 6 vector.
Library was not normalized."

ORIGIN
Query Match
Best Local Similarity 12.4%; Score 61.4; DB 13; Length 712;
Matches 53; Conservative 179; Mismatches 165; Indels 0; Gaps 0;

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/mol_type="genomic DNA"  
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_note="end : T7"
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## ORIGIN

	Query Match	10.5%; Score 51.8; DB 29; Length 683;
	Best Local Similarity	33.5%; Pred. No. 0.56;
	Matches 152; Conservative	5; Mismatches 297; Indels 0; Gaps 0;
QY	18	TATGATTGGTTGTAAATTAGACACAGGGGHTTAGGCTCTTCCATTACTTTTTTAAGTGTTTT 77
DB	485	TNTTNTTNTTNTTNTTNTTNNNAATTTTNTTNTTAAATNTTTTTTTTTTTTTTTTTTTT 426
QY	78	TCTGTTACTGTCCTCGCGAICTGATTTACGA CAATAGAGTTTCGGGTTTGTGCCCATC 137
DB	425	TTTTTTTTTTTTTGTTTTTNNNTTNTNNTNNNTTTTTTNTNTNTTNGTNNNTNTNNNN 366
QY	138	CAGTTTGGAAAATAAACGTCGCCTTTTAAAGTTTGTCTGGATCGATAACCTGTGAAGAATTG 197
DB	365	TTNTTTTTTTTTTTTTTNNNTNTTTTTTTTTTTTTTNTTNNNTNNNTTNNNTTTTTT 306
QY	198	AGCTAGTCGCAATTTATTTGGATGATCCAACTCTCATGTTTTTCTTGCTCGAAGTCT 257
DB	305	TTTTTNTNNNNTTTTTNNNTTNNNTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTNN 246
QY	258	GPTAAACAGCAATTGTCTGTCTGCGGATGTCAATACC TAGCCGTGATCGAGAACTAGG 317
DB	245	NNNTNNNNNNNTTNNNNTTGGTCTGCTNTTNNNNNNNTTTTTTNNTNTNNNTN 186
QY	318	TTTTTCGAGTCAATTTTGGCCCCCTTTTGGGTATATCTGGTTCGATTAACGATTCATCTGGAA 377
DB	185	NNNGNNNTTTTTTKTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTNTNGTGN 126
QY	378	AGGGTTTAAAGTGGTGACGTTTGTAGTATTCCAAATTTCTTCAAAATTTAGTTATGGGAATG 437
DB	125	NNTNNNNNNNTTNNNTNTTTTTTKKTTTTTWTTTTTTTTTTTTTTTTTTTTTTTATAATN 66
QY	438	AAAAATCCCGAAATGTGACTGTCAAATTCCTGTAA 471
DB	65	NBTATNTTAAATATNTTTTATTTTTTTTTTTTTTTT 32

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RESULT 8
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LOCUS
DEFINITION
CNS018GS linear GSS 26-JUL-1999
BACN13P09 of DrosBAC library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
AUI09318
GSS.
AL109318.1 GI:5629622
Drosophila melanogaster (fruit fly)
Drosophila melanogaster
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 942)
Genoscope.
Direct Submission
Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : secre@genoscope.cns.fr)
- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
http://www.edgp.ebi.ac.uk -. This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billaud at CEPH (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project grant. The DNA was prepared from embryos by Alain Bucheton
and Genevieve Pavan. It has been constructed in the vector

```

```

FEATURES             Location/Qualifiers
     source            1..942
                        /organism="Drosophila melanogaster"
                        /mol_type="genomic DNA"
                        /db_xref="taxon:7227"
                        /clone_lib="BACN13P09"
                        /clone_lib="DrosBAC"
                        /plasmid="pBelobAC11"
                        /note="end : T7"

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ORIGIN

Query Match.	10.5%;	Score 51.8;	DB 29;	Length 942;
Best Local Similarity	36.3%;	Pred. No. 0.51;		
Matches 113;	Conservative 48;	Mismatches 150;	Indels 0;	Gaps 0;
QY	159	TCITTTAAAGNTTGGCGATCGATAAACCTGCGAAGATTGAGTCGTAGTCGATTATTCGAT	218	
Db	95	TCCGWTWTTTACTTTTTTWTWTTTWWMTTWTTTWTWTTTWAATCAATAATTTGCW	154	
QY	219	GATCCCATCTTCATCGTTTTTTCTTGCTTCGAAGTTCTGTATAAACAGAAATTTGTCGTG	278	
Db	155	TWTTTWTATTTTWTWTTTWTWTTTCTKTAACGATTTTWTATAACKATWTTWTCA	214	
QY	279	TGGATTGTCATTACCTAGCGGTGATCGAGAACTAGGGTTTTCGAGTCAATTTGCCCC	338	
Db	215	WWTTTWTTCATWAAAWTTTWTWTTTWWWTKKGTTTWTWTTTWTWTTTWTWTTT	274	
QY	339	TTTTGGTTATATCTGGTTCGATAACGATTTCATCTGGATTAGGGTTTTTAAGTGGGACGTT	398	
Db	275	TTTTTTTTTWTWTTTKTGWGTRATWMTWTTTWTWTTTWTWTTTWTWTTTWTWTTT	334	
QY	399	TAGTATTTCCAATTTCTTTCAAANNTTAGTATTGGATAATGAAAATCCCGAATTCGCTGC	458	
Db	335	TKTAYTWTTTWTWTTTWTWTTTWTWTTTWTWTTTWTWTTTWTWTTTWTWTTT	394	
QY	459	AATTTCTTGTT	469	
Db	395	TWTTTWTWTTT	405	

RESULT 9	BUS34806/c
LOCUS	BUS34806
DEFINITION	AGENCOURT 10183661 NIH MGC 143 Mus musculus cDNA clone IMAGE:65452130 5', mRNA sequence.
ACCESSION	BUS34806
VERSION	BUS34806.1 GI:22845247
KEYWORDS	EST.
SOURCE	Mus musculus (house mouse)
ORGANISM	Mus musculus
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 873) NIH-MGC <a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a> . National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999)
AUTHORS	Contact: Robert Strausberg, Ph.D.
JOURNAL	Email: <a href="mailto:cgapbs-remail.nih.gov">cgapbs-remail.nih.gov</a>
COMMENT	Tissue Procurement: Dr. Michael Brownstein cDNA Library Preparation: Michael Brownstein Laboratory cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Agencourt Bioscience Corporation Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <a href="http://image.llnl.gov">http://image.llnl.gov</a> Plate: L1CM2737 row: 1 column: 02 High quality sequence stop: 141.

FEATURES	SOURCE
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/clone="IMAGE:6562130"
/lab_host="DH10B (TI-phage-resistant)"
/clone_lib="NIH_MGC_143"
/notes="Organ: Brain; Vector: pDNR-LIB; Site 1: Sfil
(gccatagggc); Site 2: Sfil (ggccgcctgggc); cDNA made
by oligo-dT priming and directionally cloned. 5' and 3'
adaptors were used in cloning as follows:
5'-AAGCAGTGATCAGCAGACAGAGGCGCATACGCGCGG-3' and
5'-ATTGAGGCGGAGCGGCGGCGCATG-3'. Full-length
enriched library was constructed using the Clontech
Creator SMART kit and size-selected to contain the 0.2-0.5
kb size fraction (other fractions present in NIH_MGC_144).
Library created in the laboratory of M. Brownstein (NIH,
NIH). Note: this is a NIH_MGC Library."

ORIGIN
Query Match 10.4%; Score 51.4; DB 13; Length 873;
Best Local Similarity 32.3%; Pred. No. 0.63;
Matches 112; Conservative 0; Mismatches 235; Indels 0; Gaps 0;

QY 23 TTTCTTTGTAATTAGATCAGGGTTAGGCTTTCCATTCTTTTAAATGTTTCTGT 82
DB TTTTCTTTTCTTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 523
QY 83 TACTGTCTCCGATCTGATTTTACGACAPAGAGTTCCGGTTTCGCCATCCAGTT 142
DB NNTTTTCTTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 463
QY 143 TGAAATPAAAGTCGCTTTTAAAGTTGCTGGATCGATAAACCTGTGAAGATGAGTCT 202
DB TTTTCTTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 403
QY 203 AGTCGATTTATGATGATCCATCTTCATCGTTTCTTCTGTCGAAAGTCTGATATA 262
DB TTTTCTTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 343
QY 263 ACCAGATTGCTGTGTCGATGTCATCTAGCGGTGATCGAGAACAGAGGTTTTC 322
DB NNNNNNTTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 283
QY 323 GAGTCAATTTGCGCCCTTTGGTTATATCTGGTTCGATAACGATTC 369
DB NNNNNNNNNNTNNNTTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 236

CNS0111D 750 bp DNA linear GSS 26-JUL-1999
Drosophila melanogaster genome survey sequence T7 end of BAC
BACN06D21 of DrosBAC library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
AL100303.1 GI:5611914
GSS.
Drosophila melanogaster (fruit fly)
Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 750)
Genoscope.
Direct Submission
Submitted (23-JUL-1999) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqrefgenoscope.cns.fr
- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of a
collaboration with the European Drosophila Genome Project (EDGP) -
http://www.edgp.ebi.ac.uk -. This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billaut at CEPH (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project Grant. The DNA was prepared from embryos by Alain Bucheton
and Genevieve Payan. It has been constructed in the vector
pBelOBAC11.
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```
Location/Qualifiers
1..750
/organism="Drosophila melanogaster"
/mol_type="genomic DNA"
/db_xref="taxon:7227"
/clone="BACN06D21"
/clone_lib="DrosBAC"
/plasmid="pBelOBAC11"
/notes="end : 17"

ORIGIN
Query Match 10.4%; Score 51.2; DB 29; Length 750;
Best Local Similarity 33.2%; Pred. No. 0.72;
Matches 113; Conservative 62; Mismatches 165; Indels 0; Gaps 0;

QY 18 TATGATTTGTTTGAATTAGATCAGGGTTTAGGCTTTCCATTCTTTTAAATGTTT 77
DB TTKTWTIKTTTTTTTAAWTTTGWKTWTGTTTTTTTWWKWTATTTTAAAGWTATK 439
QY 78 TCTGTACTGTCTCCGCGATCTGATTTTACGACATAGAGTTTCGGTTCCTCCATTC 137
DB TTTGKTTTTTCTTTTCTTCAKTKTGTGTTTTTGTGATTTTCTTTTCTTTTCTT 379
QY 138 CAGTTTGAATAAAGCGTCCTTTTAAAGTTTCTGATCGATGATAAAGCTGGAAGATG 197
DB TTTGKTTTTTCTTTTCTTCAKTKTGTGTTTTTGTGATTTTCTTTTCTTTTCTT 319
QY 198 AGTCTAGTCCGATTTATGATGATCCATTTCTTCATCGTTTTTTTCTGCTCGAAGTCT 257
DB KTAGYTTTWTGTTTGTGTTTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 259
QY 258 GTATACACGATTTGCTGTGCGATTTGCTGATGCTGCTGCTGCTGCTGCTGCTGCT 317
DB TTTGKTTTTTCTTTTCTTCAKTKTGTGTTTTTGTGATTTTCTTTTCTTTTCTT 199
QY 318 TTTTCGAGTCAATTTGCGCCCTTTGTTTATATCTGCTTC 357
DB TTTTCGAGTCAATTTGCGCCCTTTGTTTATATCTGCTTC 159

CG744146 1246 bp DNA linear GSS 24-OCT-2003
P036-3-310.ya Ppa EcoRI BAC Library Pristionchus pacificus genomic,
genomic survey sequence.
CG744146
CG744146.1 GI:37965014
GSS.
Pristionchus pacificus
Pristionchus pacificus
Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;
Neodiplogasteridae; Pristionchus.
1 (bases 1 to 1246)
Srinivasan,J., Sinz,W., Jesse,T., Wiggers-Perebolte,L., Jansen,K.,
Buntjer,J., van der Meulen,M. and Sommer,R.J.
An integrated physical and genetic map of the nematode Pristionchus
pacificus
Mol. Genet. Genomics 269 (5), 715-722 (2003)
22835951
12884007
Contact: Sommer RJ
Evolutionary Biology
Max-Planck-Institute for Developmental Biology
Spemannstr. 37-39, Tuebingen D-72076, Germany
Tel: 00497071601371
Fax: 00497071601498
Email: ralf.sommer@tuebingen.mpg.de
Class: BAC ends.
Location/Qualifiers
1..1246
/organism="Pristionchus pacificus"
/mol_type="genomic DNA"
/strain="California"

RESULT 11
CG744146
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT
FEATURES
source
```





Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the *Drosophila* melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org/TheBDGP/Drosophila>

The melanogaster BAC library was prepared by Kazutoyo Ohsesawa and Aaron Mammoler in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCT-98 and was constructed by partial *EcoRI* digestion of *Drosophila* DNA provided by the BDGP from the isogenic strain Y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at <http://bacpac.med.buffalo.edu/drosophila/bac.htm>.

**FEATURES**  
**source**

## ORIGIN

Query Match	Score 49.6;	DB 29;	Length 563;
Best Local Similarity	35.9%;	Pred. No. 1.7;	
Matches 148;	Conservative	0;	Mismatches 264;
			Indels 0;
			Gaps 0;

Qy	18	TATGATTTGTTGTAATAGATCAGGGGTAGCTCTTCCATTACTTTTAAATGTTTTT	77
Db	74	TTTTTTTTTTTTGTTTTTTTTTTTTTTTTTTGTCCTTTTTTTTTTTTTTTTTTTTT	133
Qy	78	TCGTGTACTGTCGCGGATCTGATTTACGACAATAGAGTTTCGGGTTCGTCCCAATC	137
Db	134	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTKNNNNNNNTNTNNNTNTNTNTTT	193
Qy	138	CAGTTTCGAAATAAACGTCCTCTTTAAGTTTCCTGGATCGATAAACCTGTGAAGATTG	197
Db	194	NTNNNTNNNTNTNNNTNNNTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	253
Qy	198	AGTCCTAGTCGATTTATGTGGATGACCACTTCTCATCGTTTTTTTCTGTGCTCGAAGTCT	257
Db	254	TTTTTTTTTTTTTTTTTTTTTTTTNNNNNTNNNTNTNTNTTTTTTTCTTNNNNNTNNNTN	313
Qy	258	GTATAACGAGATTTGCTGTGCGATTGTCAATACCTAGCCGTGTATCGAGAACTAGG	317
Db	314	NTNNNTNTNTTTTTTTTTTTTTTTTTTTTTTTTTNTNTNNNTNTNTNTNTNTTT	373
Qy	318	TTTTTCGAGTCAATTTTGCCCTTTTTTGTTATATCTGGTTCGATAACGATTCATCTGGATT	377
Db	374	TTTTTTTTTTTTTTTTTTTTTTTTTTTTNTNTNTNNNTNTNNNTNTNTNTNTNTNT	433
Qy	378	AGGGTTTTTAAGTGGTGCCTTTATGTATTCCAATTTCTTCCAAAATTTAGTTAT	429
Db	434	NTNTNTTTTTTTGTGNGTNNNNNTTTTTTTTTNNNNNNTTTTTTTTTTTTTTTT	485

## RESULT 14

LOCUS	CNS008BU	737 bp	DNA	linear	GSS 03-JUN-1999
DEFINITION	Drosophila melanogaster genome survey sequence TTT3 end of BAC # BACR16E03 of RPC1-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.				

AL051476	
AL051476.1	GI:4933530
Q58	
Drosophila melanogaster (fruit fly)	
Drosophila melanogaster	
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Pipera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.	
1 (bases 1 to 737)	

AUTHORS  
TITLE  
JOURNAL

COMMENT

Genoscope.  
Direct Submission  
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :  
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr  
- Web : [www.genoscope.cns.fr](http://www.genoscope.cns.fr))  
Determination of this BAC-end sequence was carried out as part of a  
collaboration with the Berkeley Drosophila Genome Project (BDGP).  
The BDGP is constructing a physical map of the Drosophila  
melanogaster genome using these BACs. For further information  
please see <http://www.fruitfly.org> The BDGP Drosophila  
melanogaster BAC library was prepared by Kazutoyo Ooeogawa and  
Aaron Mamoser in Pieter de Jong's laboratory in the Department of  
Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,  
NY. The library is named RPci-98 and was constructed by partial,  
EcoRI digestion of Drosophila DNA provided by the BDGP from the  
isogenic strain y2; cn bw sp, the same strain used for the BDGP's  
P1 and EST libraries. A more detailed description of the library  
and how to order individual BAC clones, the entire library, or  
filters for hybridization from the BACPAC Resource Center can be  
found at [http://bacpac.med.buffalo.edu/drosophila\\_bac.htm](http://bacpac.med.buffalo.edu/drosophila_bac.htm).

FEATURES	SOURCE
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## ORIGIN

Query Match	10.0%;	Score 49.6;	DB 29;	Length 737;	
Best Local Similarity	33.3%;	Pred. No. 1.5;			
Matches 138;	Conservative 14;	Mismatches 263;	Indels 0;	Gaps 0;	
18	TATGATTTGTTTGAATAGATCAGGGGTTT	AGGTCITCCATTAC	TTTTTAATGTTTTT	77	
310	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	369	
78	TCTGTTACTGTCGCGCATCTGATTTT	ACGACAATAGAGTTTCGGGTTTTGTC	CCCATTC	137	
370	NTTTTTNTNTTTNNNTTNA	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	429	
138	CAGTTTGAATAAATAACGTCGCTTTT	TAAAGTTTGTCTGGATCGATAAACCTCTG	GAAGATTG	197	
430	TTNTTTTTNTTTTTTTTTTTTTTTTTTTTT	TTTTTNTCTNTTTTTTTTTTNCANTGNNT	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	489	
198	AGTCTAGTCGATTTATGTGATCGATCCAT	CTCTTCATCGTTTTTTTTCTTGCTTC	CGAAGTTCT	257	
490	TTTTTTTNTNTTTTTNANANNNTT	MANNTNTNNTNNTNNTNNTCTTTTTNT	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	549	
258	GTATAACCGAGATTGTCGTGTGCGGATTG	TCATACCTAGCGGTGATCGAGAACTAGG	317		
550	TTTTTTTNTNTTTTTNNNNNNNNNTT	AWNNTNNSNANCNANTNNNNGNANNNNNNNN	609		
318	TTTTTCGAGTCAATTTTGCCCTTTTTGGT	TTATATCTGGTTCGATAACGATTCATCTGG	ATT	377	
610	TWTTTTTTTTTTNTNTTTTTTTTTTTTT	TTTTTTTTTTTTTTTTTTTTNNNTNTNNNT	TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	669	
378	AGGGTTTTTAAGTGGTGACGTTTAGTATTC	CAATTTCTTCAAATTTAGTTATGGA	432		
670	NNSTTTTTNNRAWNTNTNATTTTTATT	TNTNTNTTATTTNTNSATNTNTNTTNTNA	724		

RESULT 15	ACCESSION	ORGANISM
CG749401	VERSION	
LOCUS	KEYWORDS	
DEFINITION	SOURCE	

Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;  
Neodiplogasteridae; Pristionchus.

1 (bases 1 to 1403)  
Srinivasan, J., Sinz, W., Jesse, T., Wiggers-Perebolte, L., Jansen, K.,  
Buntjer, J., van der Meulen, M., and Sommer, R.J.

An integrated physical and genetic map of the nematode Pristionchus  
pacificus

Mol. Genet. Genomics 269 (5), 715-722 (2003)

22835951

12884007

Contact: Sommer RJ

Evolutionary Biology

Max-Planck-Institute for Developmental Biology

Spemannstr. 37-39, Tuebingen D-72076, Germany

Tel: 00497071601371

Fax: 00497071601498

Email: ralf.sommer@tuebingen.mpg.de

Class: BAC ends.

Location/Qualifiers

1..1403

/organism="Pristionchus pacificus"

/mol\_type="genomic DNA"

/strain="California"

/db\_xref="taxon:54126"

/clone\_lib="Ppa EcoRI BAC library"

/note="The library was generated by a partial digest of

the genomic DNA with EcoRI and cloning into the BAC

vector."

# ORIGIN

Query Match	10.0%;	Score 49.6;	DB 29;	Length 1403;
Best Local Similarity	44.2%;	Pred. No. 1.3;		
Matches 196;	Conservative	0;	Mismatches 247;	Indels 0; Gaps 0;

  

QY	27	TTTGTAAATTAGATCAGGGGTTTAGGTCCTCCATTACTTTTAAATGTTTTCGTTACT 86
Db	362	TTTGTGTTGTTT 421
QY	87	GTCTCGCGATCTGATTTTACGACATAGAGTTTCGGGTTTTGTCCCATTCAGTTGAA 146
Db	422	TTTGTGTTTGT 481
QY	147	AATAAACGTCGCTTTTAAAGTTTGTGGATCGATNAACCTGTGAAGATTGAGTCTAGTC 206
Db	482	TGTTTTTTTTTGTGTTTTTGTTTTTTTTTTTTTTTTGTGTTTTTTTGT 541
QY	207	GATTTATTCGATCATCCATCTTCATCGTTTTTTTCTTCGAAAGTTCGTATAACCA 266
Db	542	TTTTTTTTTGTGTTTTTTTGTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTGT 601
QY	267	GATTTGTCGTGCGATTGTCATTACCTAGCCGTGTATCGAGAACTAGGGTTTCGAGT 326
Db	602	TTTTTGTTTTTTTTTTTTTTTTTTTTTTTTTTTTGTGTTTTTGTGTTTTTTTGT 661
QY	327	CAATTTGCCCTTTTGGTTATCTGTTTCGATACGATTCATCTGATTAGGGTTTAA 386
Db	662	TTTTTTTGTGTTTGTGTTTTTTTTTTTTTTTGTGTTGTTGTTGTTGTTGTTT 721
QY	387	AGTGTGACGCTTAGTATTCCTCAATTTCTCAAAATTTAGTATGATGATAATGAAATCCCG 446
Db	722	GGTTTTTTTTTTTGTGTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT 781
QY	447	AATGACTGTCAATTTCTGTT 469
Db	782	TTTTTTTTTTTTTTTTTTTTTTT 804

Search completed: June 20, 2004, 05:01:22  
Job time : 2452.33 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: June 19, 2004, 23:55:13 ; Search time 366.167 Seconds

(without alignments)  
5731.297 Million cell updates/sec

Title: US-09-000-062-7

Perfect score: 494

Sequence: 1 ctcaggcgagaaacaggtat.....cgacagatcccggtatctgog 494

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 3373863 seqs, 2124099041 residues

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N\_Geneseq\_29Jan04.\*

- 1: Geneseqn1980s.\*
- 2: Geneseqn1990s.\*
- 3: Geneseqn2000s.\*
- 4: Geneseqn2001as.\*
- 5: Geneseqn2001bs.\*
- 6: Geneseqn2002s.\*
- 7: Geneseqn2003as.\*
- 8: Geneseqn2003bs.\*
- 9: Geneseqn2003cs.\*
- 10: Geneseqn2004s.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	490.8	99.4	494	2	Aat85997 Arabidops
2	374.2	75.7	1005	3	Aac47929 Arabidops
3	84.6	17.1	677	3	Aac32694 Arabidops
4	83.6	16.9	674	3	Aac51182 Arabidops
5	55.8	11.3	12870	6	Abx39983 Human che
6	55.8	11.3	12870	6	Abx39983 Human che
7	49.9	9.9	17869	6	Abx39921 Human che
8	49.9	9.9	17869	6	Abx39921 Human che
9	48.9	9.7	6593	6	Abx32478 Human inn
10	46.6	9.4	7814	4	Aas46530 Tumour su
11	46.4	9.4	6650	6	Abx32638 Human inn
12	46.2	9.4	6048	6	Abx32509 Human inn
13	46.9	9.3	3025	7	Abx210189 Haematopo
14	46.9	9.3	3025	9	Abx42119 Pretreat
15	46.9	9.3	3025	9	Abx42119 Pretreat
16	45.6	9.2	6688	6	Abx33657 Human inn
17	45.6	9.2	7442	4	Aas46686 Tumour su
18	44.8	9.1	5208	6	Abx32091 Human inn
19	44.8	9.1	12733	6	Abx32091 Human inn
20	44.4	9.0	9547	6	Abx33505 Human inn
21	44.2	8.9	594	6	Abq46990 Oligonucl
22	44.2	8.9	594	6	Abq46990 Oligonucl
23	44.4	8.9	5204	6	Abx32899 Human inn

24	44	8.9	5989	6	ABL54320	Chemical
25	44	8.9	18585	6	ABL34609	Human met
26	43.8	8.9	6467	6	ABN80169	Human che
27	43.6	8.8	18598	6	ABL32387	Human inn
28	43.4	8.8	3025	7	ABZ10043	Haematopo
29	43.4	8.8	3025	9	ADB54091	Pretreat
30	43.4	8.8	3025	9	ADB54091	Pretreat
31	43.4	8.8	6523	9	ADB84216	Human lym
32	43.4	8.8	16170	6	ABL33269	Human inn
33	43.2	8.7	3683	7	ABZ10199	Haematopo
34	43.2	8.7	3683	7	ABZ10199	Haematopo
35	43.2	8.7	8668	6	ABL33696	Human inn
36	43.2	8.7	9564	6	ABL32098	Human inn
37	43.2	8.7	12705	6	ABL32148	Human inn
38	42.8	8.7	3001	6	ABN80333	Human che
39	42.8	8.7	6120	6	ABN80333	DNA trans
40	42.8	8.7	6120	6	ABN80302	Human che
41	42.6	8.6	866	4	AAJ94068	Human neu
42	42.6	8.6	5552	6	ABL33258	Human inn
43	42.4	8.6	17918	6	AAS61418	Human inn
44	42.4	8.6	7657	4	AAS45477	Chemical
45	42.4	8.6	7657	6	ABL34022	Human inn

ALIGNMENTS

RESULT 1

AAT85997  
ID AAT85997 standard; DNA; 494 BP.

AC AAT85997;

DT 17-NOV-1997 (first entry)

XX Arabidopsis thaliana histone H3.3-like DNA fragment (intron 2).

DE Plant expression regulation sequence; intron 2; histone;  
KW herbicide tolerance; 5-enolpyruvylshikimate-3-phosphate synthase; EPSPS;  
KW glyphosate; ds.

XX Arabidopsis thaliana.

OS Arabidopsis thaliana.

PN WO9704114-A2.

PD 06-FEB-1997.

XX 17-JUL-1996; 96WO-FR001109.

PR 19-JUL-1995; 95FR-00008980.

XX (RHON ) RHONE POULENC AGROCHIMIE.

XX Derose R, Chaubet N, Gigot C;

XX WPI; 1997-132652/12.

XX New regulatory sequence for chimeric gene expression in rapidly growing parts of a plant - includes at least one intron from a plant histone gene and is useful for imparting resistance to herbicides.

PS Claim 4; Page 27; 3lpp; French.

XX The known cosmid clone c22 of Arabidopsis thaliana contains two histone H3.3-like genes. Digestion of clone c22 with restriction enzymes AluI and CfoI generated a fragment of 494 bp having the present sequence. This fragment, designated intron 2, was ligated to synthetic linkers for cloning into plant expression vectors. In addition to the intron 2 sequence, the vectors contained a plant promoter and a herbicide tolerance gene (e.g. a mutated version of the 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS) gene). The intron enhances expression of the herbicide tolerance gene in rapidly growing parts of plants. The intron can also be used to enhance expression of genes that impart resistance to

```

CC pathogens or that encode nutritional or therapeutic proteins
XX SQ Sequence 494 BP; 113 A; 83 C; 104 G; 194 T; 0 U; 0 Other;
    Query Match          99.4%; Score 490.8; DB 2; Length 494;
    Best Local Similarity 99.6%; Pred. No. 5,5e-118;
    Matches 492; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CTCAGCGAAGAACAGGTATGTTGTTGTTGAATAGATCAGGGGTTAGTCTCTTTCAT 60
DB 1 CTCAGCGAAGAACAGGTATGTTGTTGTTGAATAGATCAGGGGTTAGTCTCTTTCAT 60

QY 61 TACTTTTAAATGTTTCTGTTACTGCTCCGCGATCTGATTTTACGACAATAGAGTTT 120
DB 61 TACTTTTAAATGTTTCTGTTACTGCTCCGCGATCTGATTTTACGACAATAGAGTTT 120

QY 121 CGGGTTTCTCCATCCAGTTTGAATAAACGTCCTTTTAAAGTTTCTGCTGATCGA 180
DB 121 CGGGTTTCTCCATCCAGTTTGAATAAACGTCCTTTTAAAGTTTCTGCTGATCGA 180

QY 181 TAAACCTGTGAAGATGAGTCTAGTCGATTTTATGATGATCCATTTCTTCATCGTTTTT 240
DB 181 TAAACCTGTGAAGATGAGTCTAGTCGATTTTATGATGATCCATTTCTTCATCGTTTTT 240

QY 241 TCTTGTCTCGAAGTCTGATTAACAGATTTGTCGTGTCGATTTGATTAACCTGCGCG 300
DB 241 TCTTGTCTCGAAGTCTGATTAACAGATTTGTCGTGTCGATTTGATTAACCTGCGCG 300

QY 301 TGTATCGAAGTACAGGTTTTCGAGTCAATTTTGCCCTTTTGGTTATATCTGTTTCGAT 360
DB 301 TGTATCGAAGTACAGGTTTTCGAGTCAATTTTGCCCTTTTGGTTATATCTGTTTCGAT 360

QY 361 AACGATTCATCTGATAGGTTTAAAGTGGTGGATTTGATTTCCAAATTTCTTCAAAA 420
DB 361 AACGATTCATCTGATAGGTTTAAAGTGGTGGATTTGATTTCCAAATTTCTTCAAAA 420

QY 421 TTTAGTATGTAATGAAATCCCGAATTCGATCTTCAATTTCTGTAAATCCGCAGA 480
DB 421 TTTAGTATGTAATGAAATCCCGAATTCGATCTTCAATTTCTGTAAATCCGCAGA 480

QY 481 TCCCGGATCTCGG 494
DB 481 TCCCGGATCTCGG 494

RESULT 2
AAC47929
ID AAC47929 standard; DNA; 1005 BP.
XX XX
AC AAC47929;
XX XX
DT 18-OCT-2000 (first entry)
XX XX
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KW Hybridisation assay; Genetic mapping; Gene expression control;
KW protein identification; signal transduction pathway; metabolic pathway;
KW promoter; termination sequence; ss.
XX XX
OS Arabidopsis thaliana.
XX XX
FN EP1033405-A2.
XX XX
PD 06-SEP-2000.
XX XX
PF 25-FEB-2000; 2000EP-00301439.
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PR 29-MAR-1999; 99US-0126785P.

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KW protein identification; signal transduction pathway; metabolic pathway;  
KW Promoter; termination sequence; ss.

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 DT 18-OCT-2000 (first entry)  
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KW Hybridisation assay; genetic mapping; gene expression control;  
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 XX promoter; termination sequence; ss.  
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 OS Homo sapiens.  
 XX  
 PD WO200202806-A2.  
 XX 10-JAN-2002.  
 XX  
 PP 29-JUN-2001; 2001WO-EP007470.  
 XX  
 PP 30-JUN-2000; 2000DE-01032529.  
 XX  
 PR 01-SEP-2000; 2000DE-01043826.  
 XX  
 PA (EPIG-) EPIGENOMICS AG.  
 XX  
 XX Olek A, Piepenbrock C, Berlin K;  
 PI WPI; 2002-154757/20.  
 DR  
 XX  
 PT New nucleic acid, oligonucleotides and peptide nucleic acid-oligomers,  
 PT useful for detecting cytosine methylation state of genes associated with  
 PT pharmacogenomics and for therapy of diseases e.g. cancer.  
 XX  
 XX Claim 1; SEQ ID NO 65; 24pp; English.  
 XX  
 CC The invention relates to a nucleic acid comprising a sequence at least 18  
 CC bases in length of a segment of the chemically pretreated DNA of genes  
 CC associated with pharmacogenomics according to one of the sequences of the  
 CC genes ALDH6 (NM 000693), CYP11A (NM 000781), CYP11B (NM 000497), CYP3A3  
 CC (NM 000776 and NM 017460), DPYD (NM 000110), EPX2 (NM 001979), OCLN  
 CC (NM 002538), TXNRD1 (NM 003330), UGT8 (NM 003360), MRP (NM 004996,  
 CC NM 019901, NM 019902, NM 019862, NM 019898, NM 019899), and  
 CC their complementary sequences, or a sequence (SI) chosen from 87  
 CC sequences and their complements. The chemical pretreatment is bisulphite  
 CC treatment to convert cytosines (but not methyl-cytosines) into uracils.  
 CC Also included are an oligomer (II) in particular an oligonucleotide or a  
 CC peptide nucleic acid (PNA)-oligomer, comprising in each case at least one  
 CC base sequence having a length of 9 nucleotides which hybridises to or is  
 CC identical to a chemically pretreated DNA of genes associated with  
 CC pharmacogenomics and their complements, arranged in an array for  
 CC analysing diseases associated with the methylation state (CpG) and/or  
 CC detecting SNPs (single nucleotide polymorphisms) of the 87 sequences. The  
 CC oligomers may also be used as PCR primers. The set of 87 nucleic acids  
 CC and their complements is useful for diagnosis and therapy of solid  
 CC tumours and cancer. The present sequence represents one of the 87 DNA  
 CC sequences or its complement. Note: The sequence data for this patent did  
 CC not form part of the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
 SQ Sequence 12870 BP; 3115 A; 234 C; 3273 G; 6248 T; 0 U; 0 Other;  
 Query Match 11.3%; Score 55.8; DB 6; Length 12870;  
 Best Local Similarity 46.5%; Pred. No. 0.0002;  
 Matches 180; Conservative 0; Mismatches 207; Indels 0; Gaps 0;

QY 18 TATGATTGTTGTTTAATAGATCAGGGTTTATAGCTTTTCCATTACTTTTAAATGTTTT 77  
 DB 3147 TTTATTTGTTTAAATGTTTTTATATTTTAAAGTTTTTATGATTGATGATGTTTTT 3206  
 QY 78 TCTGTTACTGTCGCCGATCTGATTTTATGACATAGAGTTTCGGTTTGTCCATTC 137  
 DB 3207 TATAGTTTTTAAATAGATGTTTTTATGTTTATGTTTATGATAGTTTTCGTTTT 3266  
 QY 138 CAGTTTGAATAAAGTCGTCCTTTTAAAGTTTCTGGATCGATAAACCCTGGAAGATTG 197  
 DB 3267 TATTTTGTGTTTATATATGTTTTTTTTTTTGGATGTTAATTCGTTTTTTTTT 3326  
 QY 198 AGCTAGTCGATTTATGAGATGATCCATCTTCATCGTTTTTTCTTGCTTCGAGTTCT 257

DB 3327 TTTTAGTTTTTATTTGTTTTTATTTTATTTTAAATTTTATTTTATTTAGGAAGTTTT 3386  
 QY 258 GTATAACCAAGATTGCTGTGTCGCGATTGTCATTACTAGCCGCTGATCGAGAACTAGGG 317  
 DB 3387 TTTTGAATTTTATTTAGTTTATTTTATTTTATTTTATTTTGGATTTTATTTAGGAATTAAG 3446  
 QY 318 TTTTCGAGTCATTTTGGCCCTTTTGGTTATATCTCGTTGATGATAAGATTCATCTCGATT 377  
 DB 3447 TTAGTGAATACGATTGATATTTTATATATATATTTATTTGGGTTTTGGTTTTTTTTT 3506  
 QY 378 AGGTTTTTAAGTGTGTCAGCGTTTATGAT 404  
 DB 3507 TAGAAGTTCGTTAGGTATGATGATTTT 3533  
 RESULT 6  
 ABL70229  
 ID ABL70229 standard; DNA; 12870 BP.  
 XX  
 AC ABL70229;  
 XX  
 DT 01-JUL-2002 (first entry)  
 XX  
 DE Chemically treated cell signalling DNA sequence#60.  
 XX  
 KW Cell signalling; cytosine methylation; cell signalling disease; cancer;  
 KW tumour; cytostatic; ds.  
 XX  
 OS Unidentified.  
 XX  
 XX WO200202807-A2.  
 PD 10-JAN-2002.  
 XX  
 PP 29-JUN-2001; 2001WO-EP007471.  
 XX  
 PR 30-JUN-2000; 2000DE-01032529.  
 XX  
 PR 01-SEP-2000; 2000DE-01043826.  
 XX  
 PA (EPIG-) EPIGENOMICS AG.  
 XX  
 XX Olek A, Piepenbrock C, Berlin K;  
 PI WPI; 2002-154758/20.  
 DR  
 XX  
 PT Nucleic acid, useful for diagnosis and therapy of diseases associated  
 PT with cell signalling e.g. cancer, comprises chemically modified genomic  
 PT sequences of genes associated with cell signalling.  
 XX  
 XX Claim 1; SEQ ID NO 119; 24pp + Sequence Listing; English.  
 XX  
 CC The invention relates to a nucleic acid comprising a sequence of at least  
 CC 18 bases of a segment of chemically pretreated DNA of genes associated  
 CC with cell signalling. The activity of the modified sequences of the  
 CC invention may be described as cytostatic. The object of the invention is  
 CC to provide the chemically modified DNA of genes associated with cell  
 CC signalling, as well as oligonucleotides and/or PNA-oligomers for  
 CC detecting cytosine methylations, as well as a method which is  
 CC particularly suitable for the diagnosis and/or therapy of genetic and  
 CC epigenetic parameters of genes associated with cell signalling. The  
 CC chemically modified DNA provided by the invention is useful for diagnosis  
 CC and therapy of diseases such as solid tumours and cancer. The sequences  
 CC given in records ABL70111-ABL70626 represent chemically pre-treated  
 CC genomic DNA's of genes associated with cell signalling. Note: The  
 CC sequence data for this patent is not represented in the printed  
 CC specification, but is based on sequence information supplied by the  
 CC European Patent Office  
 XX  
 SQ Sequence 12870 BP; 3115 A; 234 C; 3273 G; 6248 T; 0 U; 0 Other;  
 Query Match 11.3%; Score 55.8; DB 6; Length 12870;  
 Best Local Similarity 46.5%; Pred. No. 0.0002;  
 Matches 180; Conservative 0; Mismatches 207; Indels 0; Gaps 0;



KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
KW ds.

OS Homo sapiens.

XX WO200200928-A2.

XX 03-JAN-2002.

XX 02-JUL-2001; 2001WO-EP007537.

XX 30-JUN-2000; 2000DE-01032529.

XX 01-SEP-2000; 2000DE-01043826.

XX (EPIG-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K;

XX WPI; 2002-130909/17.

XX Nucleic acid comprising fragment of chemically modified gene, useful for  
XX diagnosis and treatment of diseases associated with abnormal cytosine  
XX methylation.

XX Claim 1; SEQ ID NO 78; 32pp + Sequence Listing; German.

XX The present invention provides a number of human immune system associated  
XX genes which are modified by the methylation of cytosines. The sequences  
XX can be used in the diagnosis and treatment of immune system disorders,  
XX including eye diseases such as retinopathy, neovascular glaucoma and  
XX macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
XX leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
XX rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
XX diseases. The present sequence is a gene of the invention

XX Sequence 17869 BP; 5366 A; 158 C; 3365 G; 8978 T; 0 U; 2 Other;

Query Match 9.9%; Score 49; DB 6; Length 17869;  
Best Local Similarity 46.0%; Pred. No. 0.018;  
Matches 210; Conservative 0; Mismatches 240; Indels 7; Gaps 1;  
QY 18 TATGATTGTTGTAATAGATCAGGGTTTGGTCTTTCCATTACTTTTAAAGTTT 77  
Db 17051 TTTGTTTGTGTTTGTGAATTTATGATAGTTTATTTTGTGTTT 17110  
QY 78 TCTGTTACTGTCGCCGATCTGATTTTACGACATAGAGTTTCGGTTTGCCTATC 137  
Db 17111 TTGTTTGTGTTTATTTTCGTTTGTGTTTGTGTTTGTGTTTGTGTTT 17170  
QY 138 CAGTTTGAATAAAGCTCCGCTTTTAAAGTTTGTGATCGATCAAACTGTGAAGATTG 197  
Db 17171 GTTTTGGTTGTTTTCGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 17230  
QY 198 AGTCTAGTCGATTTATGATGATGATCATCTTCATCGTTTTCCTTCGATGTTCT 257  
Db 17231 TGT-----TTTGTGTTGTTTATTTTGTGTTGTTTGTGTTTGTGTTT 17283  
QY 258 GTATACACAGATTGCTGTCGCGATTCGATTAACCTAGCGTGTATCGACAACATG 317  
Db 17284 GGTGTTTGTGTTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 17343  
QY 318 TTTTCGAGTCAATTTTGCCTTTTCCTTTTCCTTTTCCTTTTCCTTTTCCTTT 377  
Db 17344 TTTTGTGTTTGTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 17403  
QY 378 AGGGTTTAAAGTGGTACGCTTAGTATTCATTTCTTCCAAATTTAGTTATGATG 437  
Db 17404 TTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTT 17463  
QY 438 AAAATCCGAATTCAGTGTTCATTTCTTGTAAATG 474  
Db 17464 TTTTGTGAGTTTGTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTT 17500

RESULT 9

ABL32478

XX ABL32478 standard; DNA; 6593 BP.

XX ABL32478;

XX 26-MAR-2002 (first entry)

XX Human immune system associated gene SEQ ID NO: 451.

XX Human; immune system disease; cytosine methylation; antiasthmatic;  
XX antiarteriosclerotic; antianaemic; cytosinatic; neotropic;  
XX neuroprotective; anti-HIV; anticonvulsant; ophthalmological;  
XX antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;  
XX acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
XX neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
XX ds.

XX Homo sapiens.

XX WO200200928-A2.

XX 03-JAN-2002.

XX 02-JUL-2001; 2001WO-EP007537.

XX 30-JUN-2000; 2000DE-01032529.

XX 01-SEP-2000; 2000DE-01043826.

XX (EPIG-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K;

XX WPI; 2002-130909/17.

XX Nucleic acid comprising fragment of chemically modified gene, useful for  
XX diagnosis and treatment of diseases associated with abnormal cytosine  
XX methylation.

XX Claim 1; SEQ ID NO 451; 32pp + Sequence Listing; German.

XX The present invention provides a number of human immune system associated  
XX genes which are modified by the methylation of cytosines. The sequences  
XX can be used in the diagnosis and treatment of immune system disorders,  
XX including eye diseases such as retinopathy, neovascular glaucoma and  
XX macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
XX leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
XX rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
XX diseases. The present sequence is a gene of the invention

XX Sequence 6593 BP; 1732 A; 121 C; 1448 G; 3292 T; 0 U; 0 Other;

Query Match 9.7%; Score 48; DB 6; Length 6593;

Best Local Similarity 45.8%; Pred. No. 0.018;

Matches 207; Conservative 0; Mismatches 240; Indels 5; Gaps 1;

QY 23 TTTGTTTGAATAGATCAGGGTTTGGTCTTTCCATTACTTTTAAAGTTTTCGT 82

Db 274 TTTTGTGTTTATTTATGATGATTTTATGTTTATGTTTATGTTTATGTTTATGTTT 333

QY 83 TACTGTCTCCGATCTGATTTTACGACATAGAGTTTCGGTTTTCGCCATCCAGTT 142

Db 334 TTTTGTGTTTATTTATGATGATTTTATGTTTATGTTTATGTTTATGTTTATGTTT 393

QY 143 TGAAATAAAGCTCCGCTTTTAAAGTTTGTGATCGATAAACCTGT-----GAAGATTG 197

Db 394 TTAGATTTTATTTATTTTATGTTTATTTTATTTTATTTTATTTTATTTTATTTT 453

QY 198 AGTCTAGTCGATTTATGATGATGATCCATTCATCGTTTTCCTGCTCGAAGTTCT 257

Db 454 TAAATGATTTGTTGGAGTTGATTAATTTTGTGTTGATTAAATTTGTTGTTAAATTT 513  
Qy 258 GTATACACAGATTTGCTGTGTCGCAATTTGTCATACCTAGCCGTGTATCGAGACTAGGG 317  
Db 514 TTATGATGAATTTTATTTATTTAGTATTGTAATTTTATTTTATTTAGAAATTTTATTTGGTT 573  
Qy 318 TTATTCGAGTCAATTTTGGCCCTTTTGGTTATATCTGTTTCGATAACGATTCATCTGGATT 377  
Db 574 TTTTATTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTT 633  
Qy 378 AGGTTTAAAGTGGTGACGTTTATGATTTCCAAATTTCTTCAAAATTTAGTTATGGAATG 437  
Db 634 ATTTTGTATGTTGTTTATTTGTTGTTTATTTGTTGTTTATTTGTTGTTTATTTTAAAGATGAT 693  
Qy 438 AATATCCCGAATTCACGTTTCAATTTCTTCTT 469  
Db 694 ATTTTGAATTTTGTGTAGATTAATTTATAGAT 725

RESULT 10

ABL32638  
ID AAS46530 standard; DNA; 7814 BP.  
XX AAS46530;  
AC AAS46530;  
DT 18-DEC-2001 (first entry)  
XX  
DE Tumour suppressor gene derived chemically modified sequence #252.  
XX Human; tumour suppressor gene; oncogene; antitumour; cytostatic; cancer;  
KW tumour; CpG dinucleotide; single-nucleotide polymorphism; SNP;  
KW cytosine methylation; ds.  
XX Homo sapiens.  
XX WO200168912-A2.  
XX 20-SEP-2001.  
XX 15-MAR-2001; 2001WO-EP002955.  
XX 15-MAR-2000; 2000DE-01013847.  
PR 06-APR-2000; 2000DE-01019058.  
PR 07-APR-2000; 2000DE-01019173.  
PR 30-JUN-2000; 2000DE-01032529.  
PR 01-SEP-2000; 2000DE-01043826.  
XX (EPIG-) EPIGENOMICS AG.  
XX Olek A, Piepenbrock C, Berlin K;  
PI WPI; 2001-602752/68.  
DR  
XX Fragments of chemically modified genes associated with tumor suppressor  
PT genes and oncogenes, useful in designing primers and probes for analyzing  
PT diseases associated with cytosine methylation state e.g. cancer.  
PS  
XX Claim 1; SEQ ID NO 252; 27pp; English.  
XX The invention relates to a nucleic acid comprising a sequence of 18  
CC bases, of a segment of chemically pretreated DNA (CP DNA) e.g. with  
CC bisulphite, of genes associated with tumour suppression and oncogenes  
CC having a sequence taken from 536 (actually 533 since numbers 408, 458 and  
CC 500 are missing from the sequence listing) sequences (Ss) and sequences  
CC complementary to (Ss). The nucleic acid may be a peptide nucleic acid-  
CC oligomer (PNA) of at least 9 nucleotides and may form part of a set of  
CC probes for detecting the cytosine methylation state and/or single  
CC nucleotide polymorphisms and also to be used in an array for analysing  
CC diseases associated with CpG dinucleotides e.g. cancers and tumours. The  
CC probes can also be used in a method for ascertaining genetic and/or  
CC epigenetic parameters for the diagnosis and/or therapy of existing  
CC diseases or the predisposition to specific diseases, by analysing  
CC cytosine methylations. The parameters may be compared to another set of

CC genetic and/or epigenetic parameters, the differences serving as basis  
CC for diagnosis and/or prognosis events which are disadvantageous to  
CC patients. The present sequence is one of the 533 genomic sequences  
CC derived from tumour suppressor genes and oncogenes. Sequences with even  
CC numbered seq ID numbers are the complementary sequence of the  
CC corresponding odd numbered sequence (e.g. ID 2 and ID1, ID 536 and ID  
CC 535, except for those whose partner sequence is missing). Note: The  
CC sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences

XX Sequence 7814 BP; 1677 A; 101 C; 1779 G; 4257 T; 0 U; 0 Other;

Query Match 9.4%; Score 46.6; DB 4; Length 7814;  
Best Local Similarity 46.8%; Pred. No. 0.043;  
Matches 214; Conservative 0; Mismatches 239; Indels 4; Gaps 2;

Qy 21 GATTTGTTTGTAAATTAGATCAGGGGTTTAGTCTTTCATTACTTTTAAATGTTTTTCT 80  
Db 3356 GGTTTTTTATTTAGAGAGGGTTTGGTTTTATATTTTTTTTATTTAGTTTTT 3415  
Qy 81 GTTACTGTCTCCGGATCTGA-TTTTACGACAATAGAGTTTCGGGTTTGTCCCATCCCA 139  
Db 3416 ATTATTATATTAAATTTTATAGGGTTTTTGTAGTTAGTTTTTTTATT 3475  
Qy 140 GTTGAABAATAAACGTCGCTCTTTTAACTTTGCTGATCGATAAACCTGTGAAGATTGAG 199  
Db 3476 TTTTGGTAATGTTGAAATTAATTAATTTTGGTTTATTAATTTTTTTTATT 3535  
Qy 200 TCTAGTCGATTTATGGAATCCATCTTCATCGTTTTTTCTTGTGTCGAAATTCGT 259  
Db 3536 TATTTATTTATTTGTTTTTTTATGTTATTTATTTGTTGTTTATTTTATTAGTTATT 3595  
Qy 260 ATAACAGATTTGCTGTGCGAATTCATTCACCTACCGTGATCGAGAACTAGGGTT 319  
Db 3596 ATTTATTAAGTTTATTTATGATTAATTTATGTTTATTTATTAATTTGTTGTTGTT 3655  
Qy 320 TTCGAGTCAATTTGCCCCCTTTTGGTTA---TATCTGTTTCGATAACGATTCATCTGAT 376  
Db 3656 GTTTATTTATTTGTTAGTTAAATTTATTTATTTATTTATTTATTTATTTATTTATTT 3715  
Qy 377 TAGGGTTTTTAAGTGGTGACGTTTAGTATTCCAATTTCTTCAAAATTTAGTTATGATAAT 436  
Db 3716 AATTTATTTATGTTTATTTAATTTAGTTCGTTTATTTGTTATTTAGTTATTTAGTTATTAAT 3775  
Qy 437 GAAATCCCGAATTGACTGTTCAATTTCTTGTAAAT 473  
Db 3776 TTTATTTGTTTATTTATTTATGAAATTTGTTTATTTAT 3812

RESULT 11

ABL32638  
ID ABL32638 standard; DNA; 6650 BP.  
XX  
AC ABL32638;  
XX  
DT 26-MAR-2002 (first entry)  
XX  
DE Human immune system associated gene SEQ ID NO: 611.  
XX Human; immune system disease; cytosine methylation; antiasthmatic;  
KW antiarteriosclerotic; antihaemic; cytostatic; neotropic;  
KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;  
KW antirheumatic; antiarthritic; antidiabetic; antipsoriatic;  
KW antineoplastic; cancer; eye disease; arteriosclerosis; anaemia;  
KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
XX ds.  
XX Homo sapiens.  
OS  
XX WO200200928-A2.  
XX

PD 03-JAN-2002.  
 XX 02-JUL-2001; 2001WO-EP007537.  
 PF 30-JUN-2000; 2000DE-01032529.  
 PR 01-SEP-2000; 2000DE-01043826.  
 XX (EPIG-) EPIGENOMICS AG.  
 PA Olek A, Piepenbrock C, Berlin K;  
 XX WPI; 2002-130909/17.  
 DR Nucleic acid comprising fragment of chemically modified gene, useful for  
 PT diagnosis and treatment of diseases associated with abnormal cytosine  
 PT methylation.  
 XX Claim 1; SEQ ID NO 611; 32pp + Sequence Listing; German.  
 XX The present invention provides a number of human immune system associated  
 CC genes which are modified by the methylation of cytosines. The sequences  
 CC can be used in the diagnosis and treatment of immune system disorders,  
 CC including eye diseases such as retinopathy, neovascular glaucoma and  
 CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
 CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
 CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
 CC diseases. The present sequence is a gene of the invention  
 XX Sequence 6650 BP; 1628 A; 159 C; 1561 G; 3302 T; 0 U; 0 Other;  
 SQ  
 Query Match 9.4%; Score 46.4; DB 6; Length 6650;  
 Best Local Similarity 44.0%; Pred. No. 0.047;  
 Matches 197; Conservative 0; Mismatches 251; Indels 0; Gaps 0;  
 QY 23 TTTGTTTGAATAGATCAGGGGTTAGGCTTCCATTACTTTTAAATGTTTTCTGT 82  
 DB TGTGTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTTATTTT 3657  
 QY 83 TACTGTCGCGATCGATTTTACGACATAGAGTTTCGGTTTGTCCCATTCAGTT 142  
 DB TTATTAGTTAGGTTTATAGGTTTATATATAAGTGTTCGGTATTGTTGATTTATGA 3717  
 QY 143 TGAATTAACGTCGTTTAAAGTTTCTGTCGATCGATAAACCTGTGAAGTTGAGTCT 202  
 DB TGATTTTAAGATGTTTGTGTAAGTTTAAATTTGTTAATTTATTTATTTAATAAT 3777  
 QY 203 AGTCGATTTATGGATCATCCATCTCTCAATCGTTTTTTTCTTCGTCGAGTTCTGTATA 262  
 DB ACCGAATGACAGTGGTAGTGATTAATGAGTATATATGTTTTTTTGTATGTTATTTTC 3837  
 QY 263 ACCAGATTGCTGTGCGGATTTGATACCTAGCCGTATCGAGAACTAGGGTTTTC 322  
 DB GTTTTTTATAATTTAGATTAGTAATTTTGTGTAATTTGATTTTGAATTTATTTT 3897  
 QY 323 GAGTCAATTTGCCCCTTTGGTTATATCTGTTTCCGATACGATTCATCTGGAATAGGGT 382  
 DB TATTTTATTTTATGGGTTTATGTTTGTGTTTGTATTTTATTTTATTTTGAAT 3957  
 QY 383 TTTAAGTGGTGACGTTTAGTATTCCTCAAAATTTAGTATGGAATGAAT 442  
 DB TCGTTTTTGAATGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTTGTT 4017  
 QY 443 CCCGAATGACTGTTCAATTTCTTGTTA 470  
 DB ATTTGGTTGATTTGTTGAATTTTATTTT 4045  
 RESULT 12  
 ID ABL32509  
 AC ABL32509 standard; DNA; 6048 BP.  
 XX ABL32509;  
 XX

DT 26-MAR-2002 (first entry)  
 XX Human immune system associated gene SEQ ID NO: 482.  
 DE Human; immune system disease; cytosine methylation; antiasthmatic;  
 XX neuroproliferative; anti-HIV; anticonvulsant; ophthalmological;  
 KW antiarthritic; antidiabetic; antipsoriasis;  
 KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;  
 KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;  
 KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;  
 KW ds.  
 XX Homo sapiens.  
 OS WO200200928-A2.  
 XX 03-JAN-2002.  
 PD 02-JUL-2001; 2001WO-EP007537.  
 PF 30-JUN-2000; 2000DE-01032529.  
 PR 01-SEP-2000; 2000DE-01043826.  
 XX (EPIG-) EPIGENOMICS AG.  
 PA Olek A, Piepenbrock C, Berlin K;  
 XX WPI; 2002-130909/17.  
 DR Nucleic acid comprising fragment of chemically modified gene, useful for  
 PT diagnosis and treatment of diseases associated with abnormal cytosine  
 PT methylation.  
 XX Claim 1; SEQ ID NO 482; 32pp + Sequence Listing; German.  
 XX The present invention provides a number of human immune system associated  
 CC genes which are modified by the methylation of cytosines. The sequences  
 CC can be used in the diagnosis and treatment of immune system disorders,  
 CC including eye diseases such as retinopathy, neovascular glaucoma and  
 CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid  
 CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,  
 CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel  
 CC diseases. The present sequence is a gene of the invention  
 XX Sequence 6048 BP; 1296 A; 103 C; 1361 G; 3288 T; 0 U; 0 Other;  
 SQ  
 Query Match 9.4%; Score 46.2; DB 6; Length 6048;  
 Best Local Similarity 46.9%; Pred. No. 0.051;  
 Matches 144; Conservative 0; Mismatches 163; Indels 0; Gaps 0;  
 QY 163 TTAAGTTTGTGGATCGATAAACCTGTGAAGATTGAGTCTAGTCTAGTTTATTTGATGATC 222  
 DB TTAAGTTTGTGGATCGTTTAAAGTTTATTAAGTTTATTAAGTTTATTTTGT 2384  
 QY 223 CATCTTCATCGTTTTTTTCTGCTCGAAGTTCTGTATTAACAGATTGCTGTGCG 282  
 DB GATTTTTTTTTGTTATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGT 2444  
 QY 283 ATTGTCATTACTAGCCGTGTATCGAGAACTAGGTTTTCGAGTCAATTTTCCCCCTTT 342  
 DB TTTTGAATTTGTTAGTTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGT 2504  
 QY 343 GGTATATCTGTTTCGATACGATTCATCTCGATTAGGTTTAAAGTGTGACGTTAGT 402  
 DB TTTGATTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGT 2564  
 QY 403 ATTCGAATTTCTCAAAATTTAGTTTATGTAATCGAAATCCGAATTTGACTGTCAAT 462  
 DB GTTTAAATGTTATTTTATTTAGTTTGTGAGTTTGTGAGTTTGTGAGTTTGTGAGT 2624  
 QY 463 TCTTGT 469  
 |||||

27-FEB-2003; 2003WO-EP002035.  
27-FEB-2002; 2002EP-00004551.  
(EPIG-) EPIGENOMICS AG.  
Adorjan P, Burger M, Maier S, Nimmrich I, Becker E, Lesche R;  
Rujan T, Schmitt A;  
WPI; 2003-731620/69.  
Detecting and differentiating between colon cell proliferative disorders  
associated with a gene or its regulatory regions comprises contacting a  
target nucleic acid in a biological sample obtained from the subject with  
a reagent.  
Claim 32; SEQ ID NO 275; 74pp; English.  
The invention relates to a novel method for detecting and differentiating  
between colon cell proliferative disorders associated with at least one  
gene or its regulatory regions. The method comprises contacting a target  
nucleic acid in a biological sample obtained from the subject with at  
least one reagent or a series of reagents, where the reagent or series of  
reagents, distinguishes between methylated and non methylated CpG  
dinucleotides within the target nucleic acid. The molecules of the  
invention demonstrate cytotatic activity whilst the method may be useful  
for detecting and differentiating between colon cell proliferative  
disorders, including cancers such as colon adenoma and colon carcinoma.  
The FNA (peptide nucleic acid)-oligomers are useful as probes for  
determining cytosine methylation state or single nucleotide  
polymorphisms. The current sequence is that of the pretreated genomic DNA

2625 TTTTTH 2631

RESULT 13

ABZ10189

ABZ10189 standard; DNA; 3025 BP.

ABZ10189;

16-JAN-2003 (first entry)

Haematopoietic cell proliferation disorder related DNA sequence #329.

Human; haematopoietic cell proliferation disorder; cytostatic; gene therapy; lymphocytic leukaemia; acute myelogenous leukaemia; cytosine methylation state; gene; ds.

Homo sapiens.

WO200277272-A2.

03-OCT-2002.

26-MAR-2002; 2002WO-EP003401.

26-MAR-2001; 2001US-0278333P.

(EPIG-) EPIGENOMICS AG.

Berlin K, Braun A, Distler J, Guetig D, Howe A, Mueller J; Olek A, Piepenbrock C, Adorjan P, Grabs G, Lesche R, Leu E; Lewin A, Lipscher E, Maier S, Model F, Mueller V, Otto T, Pelet C; Schwöbe I, Ziebarth H;

WPI; 2003-018942/01.

Detecting and differentiating between hematopoietic cell proliferative disorders, comprises contacting a target nucleic acid with a reagent that distinguishes between methylated and non-methylated CpG dinucleotides.

Claim 28; SEQ ID NO 329; 117pp; English.

The present invention describes a method for detecting and differentiating between haematopoietic cell proliferative disorders associated with at least 1 gene and/or their regulatory regions in a subject. The method comprises contacting a target nucleic acid in a biological sample obtained from the subject with at least 1 reagent, which distinguishes between methylated and non-methylated CpG dinucleotides within the target nucleic acid. ABZ09861 to ABZ1118 represent specifically claimed nucleotide sequences from the present invention. Oligonucleotides from the present invention can be used for differentiating between healthy haematopoietic cells and proliferative disorder haematopoietic cells; for differentiating between acute lymphocytic leukaemia and acute myelogenous leukaemia; as probes for determining the cytosine methylation state and/or single nucleotide polymorphisms (SNPs) of haematopoietic cell proliferation disorder related sequences and their complements; and as primers for the amplification of haematopoietic cell proliferation disorder related DNA sequences. The nucleotide sequences from the present invention can also be used for detecting a predisposition to, differentiation between subclasses, diagnosis, prognosis, treatment and/or monitoring of haematopoietic cell proliferative disorders. The present method enables a highly specific classification of haematopoietic cell proliferative disorders allowing for improved and informed treatment of patients

Sequence 3025 BP; 535 A; 0 C; 930 G; 1560 T; 0 U; 0 Other;

Query Match 9.3%; Score 46; DB 7; Length 3025;

Best Local Similarity 47.9%; Pred. No. 0.049;

Matches 162; Conservative 0; Mismatches 175; Indels 1; Gaps 1;

23 TTTGTTTGTAAATAGATCAGGGGTAGGCTTTTCCTACTTTTAAATGTTTTCGT 82



GenCore version 5.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: June 20, 2004, 00:55:23 ; Search time 3463.42 Seconds  
(without alignments)  
6182.178 Million cell updates/sec

Title: US-09-000-062-7  
Perfect score: 494  
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Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 3470272 seqs, 21671516995 residues

Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0  
Maximum DB seq length: 20000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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- 2: gb\_hgt.\*
- 3: gb\_in.\*
- 4: gb\_on.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sv.\*
- 13: gb\_un.\*
- 14: gb\_vi.\*
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- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
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- 22: em\_ov.\*
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- 31: em\_hgt\_inv.\*
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- 33: em\_hgt\_mus.\*
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- 35: em\_hgt\_rdt.\*
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- 37: em\_hgt\_vrt.\*
- 38: em\_sy.\*
- 39: em\_hgt\_hum.\*
- 40: em\_hgt\_mus.\*
- 41: em\_hgt\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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C 5	477.8	96.7	118267	8	ATTSJ17	AD035708 Arabidops
6	476.2	96.4	777	8	AJ592632	AJ592632 Arabidops
7	55.8	11.3	12870	6	AX348370	AX348370 Sequence
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C 9	50.6	10.2	14281	8	SKORFS	X54850 S.klyveri
10	49.6	10.0	7218	6	I66494	I66494 Sequence 14
11	49.6	10.0	34980	6	AX344573	AX344573 Sequence
12	49	9.9	17869	6	AX345007	AX345007 Sequence
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C 14	49	9.9	176162	5	AL953898	AL953898 Zebrafish
15	48.6	9.8	250029	3	A2014820	AE014820 Plasmodi
16	48	9.7	6593	6	AX345380	AX345380 Sequence
17	47.8	9.7	93791	2	AC138073	AC138073 Homo sapi
C 18	47.2	9.6	165888	2	AC144291	AC144291 Macaca mu
C 19	47.2	9.6	166224	2	AC142946	AC142946 Macaca mu
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21	47	9.5	95764	2	AC015732	AC015732 Homo sapi
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39	45.6	9.2	170827	2	AC125567	AC125567 Rattus no
40	45.6	9.2	236566	2	AC094427	AC094427 Rattus no
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ALIGNMENTS

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VERSION	AR182672.1	GI:20225879				
KEYWORDS	Unknown.					
SOURCE	Unknown.					
ORGANISM	Unclassified.					
REFERENCE	1 (bases 1 to 494)					
AUTHORS	Derose, R., Chaubet, N. and Gigot, C.					
TITLE	Isolated DNA sequence capable of serving as regulatory element in a chimeric gene which can be used for the transformation of plants					
JOURNAL	Patent: US 6338961-A 7 15-JAN-2002;					



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  Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
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  Title
  Journal
  Comment
  Other publication FR 2736929 970124.
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  Version
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  Source
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  Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
  rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
  1 (bases 1 to 4833)
  Reference
  Authors
  Title
  Journal
  Comment
  Other publication
  Location/Qualifiers
    J. Mol. Biol. 225 (2), 569-574 (1992)
    92277653
    1593639
    2 (bases 1 to 4833)
    Gigot,C.
    Direct Submission
    Submitted (17-SEP-1991) C. Gigot, Inst de Biologie Mol des Plantes,
    12 Rue du General Zimmer, 67084 Strasbourg Cedex, FRANCE
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ACCESSION AL161596
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REFERENCE	1 (bases 1 to 107700) Hudson, S., Mewes, H.W., Lemcke, K. and Murphy, G., Ridley, P.,		
AUTHORS	Murphy, G., Ridley, P., Hudson, S., Mewes, H.W., Lemcke, K. and Mayer, K.F.X.		
JOURNAL	Unpublished		
REFERENCE	2 (bases 9546 to 9977) Volckaert, G., Grymonprez, B., Voet, M., Robben, J., Mewes, H.W., Lemcke, K. and Mayer, K.F.X.		
AUTHORS	Unpublished		
JOURNAL	Unpublished		
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AUTHORS	Unpublished		
JOURNAL	Unpublished		
REFERENCE	4 (bases 1 to 112067) EU Arabidopsis sequencing, project.		
AUTHORS	Direct Submission		
TITLE	Submitted (10-MAR-2000) MIPS, at the Max-Planck-Institut fuer Biochemie, Am Klopferspitz 18a, D-82152 Martinsried, FRG, E-mail: lemcke@mips.biochem.mpg.de, mayer@mips.biochem.mpg.de Project		
JOURNAL	Coordinator: Mike Bevan, Molecular Genetics Department, Cambridge Laboratory, John Innes Centre, Colney Lane, NR4 7UJ Norwich, UK, E-mail: michael.bevan@bbsrc.ac.uk		
COMMENT	Information on performance of analysis and a more detailed annotation of this entry and other sequences of chromosomes 3, 4 and 5 can be viewed at: <a href="http://www.mips.biochem.mpg.de/proj/thal/">http://www.mips.biochem.mpg.de/proj/thal/</a> and this fragment has an overlap with ATCHRIV91 at the 5' end and an overlap with ATCHRIV93 at the 3' end.		
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Matches 479; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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QY 301 TGTATCGAAGACTAGGTTTTTCGAGTCAATTTGCCCCCTTTTGGTTATATCTGGTTCCAT 360

DB 83615 TGTATCGAAGACTAGGTTTTTCGAGTCAATTTGCCCCCTTTTGGTTATATCTGGTTCCAT 83556

QY 361 AACGATTTCATCTCGATTAGGGTTTTTAAGTGGTGAGCTTTAGTATTCCCAATTTCTTCAAAA 420

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DB 83495 TTTAGTTATGGAATAAGAAATCCGAATGACCTGTTCAATTTCTTGTAAATGCGCAGA 83436

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DB 83435 T 83435

RESULT 5  
LOCUS ATT5J17/c 118267 bp DNA linear PLN 21-SEP-1999  
DEFINITION Arabidopsis thaliana DNA chromosome 4, BAC clone T5J17 (ESSA project).

ACCESSION AL035708  
VERSION AL035708.2 GI:5918309

KEYWORDS Arabidopsis thaliana (thale cress)

SOURCE Arabidopsis thaliana

ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1  
AUTHORS Bevan, M., Murphy, G., Ridley, P., Hudson, S., Bancroft, I., Mewes, H.W., Mayer, K.F.X., Lemcke, K. and Schueller, C.  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 118267)  
AUTHORS EU Arabidopsis sequencing, project.  
TITLE Direct Submission  
JOURNAL Submitted (21-SEP-1999) MIPS, at the Max-Planck-Institut fuer Biochemie, Am Klopferspitz 18a, D-82152 Martinsried, FRG, E-mail: schueller@mips.biochem.mpg.de, mayer@mips.biochem.mpg.de, Project Coordinator: Mike Bevan, Molecular Genetics Department, Cambridge Laboratory, John Innes Centre, Colney Lane, NR4 7UU Norwich, UK, E-mail: michael.bevan@bbsrc.ac.uk

COMMENT On Sep 22, 1999 this sequence version replaced gi:4490734. Information on performance of analysis and a more detailed annotation of this entry and other sequences of chromosomes 3, 4 and 5 can be viewed at: <http://www.mips.biochem.mpg.de/proj/thal/>.

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QY 481 T 481
DB 94002 T 94002

RESULT 6
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LOCUS       Arabidopsis thaliana T-DNA flanking sequence, right border, clone
DEFINITION 627A08.
ACCESSION  AJ592632
VERSION     AJ592632.1 GI:37942256
KEYWORDS   right border; T-DNA flanking sequence.
SOURCE     Arabidopsis thaliana (Chale cress)
ORGANISM   Arabidopsis thaliana
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            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE
1
  Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
  Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
  Lepiniec, L., Caboche, M. and Lecharny, A.
  T-DNA integration into the Arabidopsis genome depends on sequences
  of pre-insertion sites
  EMBO Rep. 3 (12), 1152-1157 (2002)

JOURNAL    EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE    22363535
PUBMED     12446585
REFERENCE  2 (bases 1 to 777)
AUTHORS    Balzerque, S.
TITLE      Submitted (23-OCT-2003) Balzerque S., UMRGV, INRA/CNRS, 2 rue
JOURNAL    Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT    PCR was performed on DNA from transformants of Arabidopsis thaliana

plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).

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Best Local Similarity 99.4%; Pred. No. 4.3e-101;
Matches 478; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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QY 61 TACTTTTTTAAGTTTTTCTGTTACTGTCTCCGGATCTGATTTTACGACAATAGAGTTT 120
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QY 481 T 481
DB 617 T 617

RESULT 7
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DEFINITION 65 from Patent WO0202806.
ACCESSION  AX348370
VERSION     AX348370.1 GI:18614406
KEYWORDS   synthetic construct
SOURCE     synthetic construct
ORGANISM   synthetic construct
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CDS

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CDS

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Query Match 10.2%; Score 50.6; DB 8; Length 14281;  
Best Local Similarity 48.8%; Pred. No. 0.099;  
Matches 137; Conservative 0; Mismatches 144; Indels 0; Gaps 0;

QY 195 TTGAGCTAGTCGATTTATGGATGATCCATCTTCATCGTTTTTTCTTCTGTCGACGT 254  
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QY 255 TCTGTATAACACGATTTGCTGTGCGATGTGCTATACCTAGCCGTGATCGAGACTA 314  
DB 11230 TTGATCTATATAAGTTTTATGTTTCGTTATAGTTACTTGACCTATTTTTCGAACAT 11171

QY 315 GGGTTTCGAGTCAATTTGCCCTTTTGGTTATATCTGCTCGATAACGATTCATCTGG 374  
DB 11170 ATGCTATAATTAATCTTTGCTTTAGATCTCTTTTCATCATTAATTTATGCTTTG 11111

QY 375 ATTAGGGTTTTTAAGTGTGTCAGCTTTAGTATTCCTCAATTTCTCAAAATTTAGTTATGATA 434  
DB 11110 TCTATTTGGGTCAATCTTTATTTATTTATTTCTTTCTCAATATATAGTCTTGATTT 11051

QY 435 ATGMAATCCGAAATGACGTGTTCAATTTCTGTTAAATCC 475  
DB 11050 ATCTTCTTAACTAATTAATTTTGTTTTAAATCTGTATAC 11010

RESULT 10  
LOCUS I66494 7218 bp DNA linear PAT 28-DEC-1997  
DEFINITION Sequence 14 from patent US 5670367.  
ACCESSION I66494  
VERSION I66494.1 GI:2724471  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (Bases 1 to 7218)  
AUTHORS Dorner, F., Scheifflinger, F. and Falkner, F. Gunter.  
TITLE Recombinant fowlpox virus  
JOURNAL Patent: US 5670367-A 14 23-SEP-1997;  
FEATURES Location/Qualifiers  
source 1..7218  
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Best Local Similarity 6.1%; Pred. No. 0.18;  
Matches 25; Conservative 213; Mismatches 172; Indels 0; Gaps 0;

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DB 54643 GTTTTITAGTAAAGTTTTTTTGTGTTTAAATAAAGGAGATTAAGAGTACG 54702

Db 1166 YY 1225

QY 194 ATTGAGCTTAGTCGATTTATGGATGATCCATCTCTTCATCGTTTTTTCTGTTTTCGAAG 253  
DB 1226 YY 1285

QY 254 TTTCTGTATACACGATTTGCTGTGCGATTTGCTATTACTACCGCTGATCGAGAAT 313  
DB 1286 YY 1345

QY 314 AGGGTTTTCGAGTCAATTTTCCCTTTTGGTTATATCTGCTGATCGATAACGATTCATCTG 373  
DB 1346 YY 1405

QY 374 GATTAGGGTTTAAAGTGGTGGCTTTAGTATTCATTCATCTTCTCAAAATTT 423  
DB 1406 YYYGGTACCAATTTCTTATCTCT 1455

RESULT 11  
LOCUS AX344573 349980 bp DNA linear PAT 01-FEB-2002  
DEFINITION Sequence 24 from Patent WO0200932.  
ACCESSION AX344573  
VERSION AX344573.1 GI:18492459  
KEYWORDS synthetic construct  
SOURCE synthetic construct  
ORGANISM artificial sequences.  
REFERENCE 1  
AUTHORS Olek, A., Piepenbrock, C. and Berlin, K.  
TITLE Diagnosis of known genetic parameters within the mhc  
JOURNAL Patent: WO 0200932-A 24 03-JAN-2002;  
Epi-genomics AG (DE)  
FEATURES Location/Qualifiers  
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Query Match 10.0%; Score 49.6; DB 6; Length 349980;  
Best Local Similarity 49.6%; Pred. No. 0.13;  
Matches 127; Conservative 0; Mismatches 129; Indels 0; Gaps 0;

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QY 78 TCTGTACTGTCTCGCATCTGATTTTACGACATAGATTTTCGGTTTTCCTCCATC 137  
DB 54583 TTTTGTAGTAATTTTATATTTTTTTTTTAAAGTTTTTTTAAAGAAATTTTATTTT 54642

QY 138 CAGTTTGAATAAAGCTCCGCTTTTAAAGTTTGTGATCGATAAACCTGTGAAGATTG 197  
DB 54643 GTTTTITAGTAAAGTTTTTTTGTGTTTAAATAAAGGAGATTAAGAGTACG 54702







Qy 403 ATTCCAAATTCCTCAAAAATTAGTTATGGATAATGAAAAATCCGAAATTCGACTGTTCATTT 462  
Db 129446 TTKRTWWTYYTTTWWTTTKKTAAWAAWKKWWWWYCWAAWTTTGRWKTCTCMMMAAAGKT 129505  
Qy 463 TCTTGTTAAATGGCGAGATCC 483  
Db 129506 TYCMRCTAWTTTWWYATCTTC 129526

Search completed: June 20, 2004, 03:45:49  
Job time : 3467.42 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: June 20, 2004, 01:40:43 ; Search time 59.125 Seconds  
(without alignments)  
3923.374 Million cell updates/sec

Title: US-09-000-062-6  
Perfect score: 418  
Sequence: 1 ttaggtacgattcttcgac.....taatttggtaacagatccc 418

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 682709 seqs, 277475446 residues

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued Patents NA.\*  
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2: /cgn2\_6/ptodata/2/ina/5B COMB.seq.\*  
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5: /cgn2\_6/ptodata/2/ina/PTCUS COMB.seq.\*  
6: /cgn2\_6/ptodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	418	100.0	418	4	US-09-000-062-6
2	43.8	10.5	5852	1	Sequence 6, Appli
3	42.4	10.1	8537	4	Sequence 2, Appli
4	40.2	9.6	5455	4	Sequence 41, Appl
5	40.2	9.6	11049	4	Sequence 34, Appl
6	39.6	9.5	11015	4	Sequence 23, Appl
7	39.4	9.4	5340	4	Sequence 55, Appl
8	39.2	9.4	11050	4	Sequence 21, Appl
9	39	9.3	5501	4	Sequence 86, Appl
10	38.4	9.2	5562	4	Sequence 37, Appl
11	38.2	9.1	5501	4	Sequence 63, Appl
12	38.2	9.1	6801	4	Sequence 38, Appl
13	38.2	9.1	6886	4	Sequence 61, Appl
14	38	9.1	786431	4	Sequence 20, Appl
15	37.8	9.0	5455	4	Sequence 3, Appli
16	37.8	9.0	6801	4	Sequence 33, Appl
17	37.8	9.0	19124	2	Sequence 62, Appl
18	37.6	9.0	11050	4	Sequence 13, Appl
19	37.6	9.0	11811	3	Sequence 85, Appl
20	37.4	8.9	5666	4	Sequence 7, Appli
21	37.4	8.9	6306	4	Sequence 29, Appl
22	37.4	8.9	23439	4	Sequence 50, Appl
23	37.2	8.9	832	4	Sequence 38, Appl
24	37	8.9	6317	4	Sequence 2813, Ap
25	37	8.9	10640	4	Sequence 11, Appl
26	36.6	8.8	984	4	Sequence 5, Appli
27	36.6	8.8	19513	4	Sequence 2100, Ap
					Sequence 40, Appl

c	28	36.6	8.8	640681	4	US-09-790-988-1	Sequence 1, Appli
	29	36.4	8.7	6040	4	US-10-204-708-70	Sequence 70, Appl
	30	36.4	8.7	6182	4	US-10-204-708-88	Sequence 88, Appl
	31	36.4	8.7	8607	4	US-10-204-708-71	Sequence 71, Appl
	32	36.4	8.7	202001	4	US-09-734-674-3	Sequence 3, Appli
	33	36.2	8.7	6113	4	US-10-204-708-14	Sequence 14, Appl
	34	36.2	8.7	7218	1	US-08-232-463-14	Sequence 14, Appl
	35	36.2	8.7	8093	4	US-10-204-708-32	Sequence 32, Appl
	36	36.2	8.7	11015	4	US-10-204-708-56	Sequence 56, Appl
	37	36	8.6	6070	4	US-10-204-708-10	Sequence 10, Appl
	38	36	8.6	19233	4	US-10-204-708-45	Sequence 45, Appl
c	39	35.8	8.6	2103	4	US-09-543-681A-797	Sequence 797, Appl
	40	35.8	8.6	5666	4	US-10-204-708-30	Sequence 30, Appl
	41	35.8	8.6	8607	4	US-10-204-708-72	Sequence 72, Appl
	42	35.4	8.5	522	4	US-09-134-001C-1017	Sequence 1017, Ap
	43	35.4	8.5	8961	4	US-10-204-708-80	Sequence 80, Appl
c	44	35.2	8.4	6243	2	US-09-056-075-1	Sequence 1, Appli
	45	35.2	8.4	8537	4	US-10-204-708-42	Sequence 42, Appl

ALIGNMENTS

RESULT 1  
US-09-000-062-6  
; Sequence 6, Application US/09000062  
; Patent No. 6338961  
; GENERAL INFORMATION:  
; APPLICANT: DEROSE, Richard  
; APPLICANT: CHAUBET, Nicole  
; TITLE OF INVENTION: ISOLATED DNA SEQUENCE CAPABLE OF SERVING AS REGULATORY  
; TITLE OF INVENTION: ELEMENT IN A CHIMERIC GENE WHICH CAN BE USED FOR THE  
; TITLE OF INVENTION: TRANSFORMATION OF PLANTS  
; FILE REFERENCE: 022650-453  
; CURRENT APPLICATION NUMBER: US/09/000,062  
; CURRENT FILING DATE: 1998-05-29  
; EARLIER APPLICATION NUMBER: PCT/FR96/01109  
; EARLIER FILING DATE: 1996-07-17  
; EARLIER APPLICATION NUMBER: FR 95/08980  
; EARLIER FILING DATE: 1995-07-19  
; NUMBER OF SEQ ID NOS: 22  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 6  
; LENGTH: 418  
; TYPE: DNA  
; ORGANISM: Zea mays  
US-09-000-062-6

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Matches 418;	Conservative 0;			
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Db	1	1	1	1
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Qy	121	1	1	1
Db	121	1	1	1
Qy	181	1	1	1
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Qy	241	1	1	1
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Db	3003	GTTCCTTTGCTTTTATTATTATTGTTATCGATTTTATAAGTTTTTAAAGAGAAAGTT	3062							
Qy	269	CTTATTGCGGCGATTGTTGATTAGGGTTTTGATTTCTTGATTATGCGATTGCAATTAGGG	328							
Db	3063	TGTTATATGTTGTTGTTTATAGAAAGTGTTTTATTAAATTGATTGGTTAGGAGGTAAATAG	3122							
Qy	329	ATTTCTCTCTGTTTTGCGTTGA	350							
Db	3123	GTTCCTTTTCAATATGTGTTTA	3144							

RESULT 7  
US-09-627-122-21  
; Sequence 21, Application US/09627122  
; Patent No. 6472521  
; GENERAL INFORMATION:  
; APPLICANT: Uhlmann, Eugen  
; APPLICANT: Greiner, Beate  
; APPLICANT: Unger, Eberhard  
; APPLICANT: Gothe, Gislinde  
; APPLICANT: Schwarzel, Marc  
; TITLE OF INVENTION: OLIGONUCLEOTIDES FOR THE INHIBITION OF HUMAN eg5  
; FILE REFERENCE: 02481.1678  
; CURRENT APPLICATION NUMBER: US/09/627,122  
; CURRENT FILING DATE: 2000-07-27  
; NUMBER OF SEQ ID NOS: 23  
; SOFTWARE: Patentin Ver. 2.1  
; SEQ ID NO 21  
; LENGTH: 5340  
; TYPE: DNA  
; ORGANISM: Plasmodium falciparum  
US-09-627-122-21

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RESULT 8
US-10-204-708-86
; Sequence 86, Application US/10204708
; Patent No. 6577731
; GENERAL INFORMATION:
; APPLICANT: OLEX, Alexander
; APPLICANT: PIEPENBROCK, Christian
; APPLICANT: BEBLIN, Kurt
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
; TITLE OF INVENTION: by Assessing DNA Methylation

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/ FILE REFERENCE: 5013.1012
/ CURRENT APPLICATION NUMBER: US/10/204,708
/ CURRENT FILING DATE: 2003-05-06
/ PRIOR APPLICATION NUMBER: PCT/EP01/03971
/ PRIOR FILING DATE: 2001-04-06
/ PRIOR APPLICATION NUMBER: DE 10019058.8
/ PRIOR FILING DATE: 2000-04-06
/ PRIOR APPLICATION NUMBER: DE 10019173.8
/ PRIOR FILING DATE: 2000-04-07
/ PRIOR APPLICATION NUMBER: DE 10032529.7
/ PRIOR FILING DATE: 2000-06-30
/ PRIOR APPLICATION NUMBER: DE 10043826.1
/ PRIOR FILING DATE: 2000-09-01
/ NUMBER OF SEQ ID NOS: 98
/ SEQ ID NO 86
/ LENGTH: 11050
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-204-708-86

Query Match          9.4%; Score 39.2; DB 4; Length 11050;
Best Local Similarity 46.9%; Pred. No. 0.27;
Matches 122; Conservative 0; Mismatches 138; Indels 0; Gaps 0

QY      154  GTATAAGATTTTAGGTTAGATGATCGTATAGCTAGATTTCATCACCAGATGTTTC 213
Db      268  GTAATATTCGATAAATGATGATTTTTTTTATTGTTATTAGTATTTTGATATGGGTTG 327

QY      214  TTGTCTAGAAATCTCGAAATCTTCGATAGTTTTCACATGCTGTAATAGATGTTCTTAT 273
Db      328  TATGACGTGGGTTTGGTAGATTTTTTTTATTATTTCGTTATATTTTATCGTTTTT 387

QY      274  TCGGCGATGTTGATTAGGTTTTGATTTCTTGATTATCGGATTGCAATTAGCGATTTT 333
Db      308  TGAGTTTTTAATTATATGTTTTAGTTTGATATTAATTTTGTATTAGGGTTTTTG 447

QY      334  CTTTGGTTTTGTGTGATCTTACGATACATCTCGCAATTGAATAGATGATCTAAAT 393
Db      448  AATTGGTGTGTTTTTTTATTGGAAATTTTTTTTTTTTGAGGTTATATGCTGGTTTTT 507

QY      394  CTTGTTAAATTTGTTCAACAG 413
Db      508  TTAGTTATTTAGTTTTAGAG 527

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RESULT 9  
US-10-204-708-37  
/ Sequence 37, Application US/10204708  
/ Patent No. 6677731  
/ GENERAL INFORMATION:  
/ APPLICANT: OLEK, Alexander  
/ APPLICANT: PIEPENBROCK, Christian  
/ APPLICANT: BERLIN, Kurt  
/ TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication  
/ TITLE OF INVENTION: by Assessing DNA Methylation  
/ FILE REFERENCE: 5013.1012  
/ CURRENT APPLICATION NUMBER: US/10/204,708  
/ CURRENT FILING DATE: 2003-05-06  
/ PRIORITY APPLICATION NUMBER: PCT/EP01/03971  
/ PRIOR FILING DATE: 2001-04-06  
/ PRIOR APPLICATION NUMBER: DE 10019058.8  
/ PRIOR FILING DATE: 2000-04-06  
/ PRIOR APPLICATION NUMBER: DE 10019173.8  
/ PRIOR FILING DATE: 2000-04-07  
/ PRIOR APPLICATION NUMBER: DE 10032529.7  
/ PRIOR FILING DATE: 2000-06-30  
/ PRIOR APPLICATION NUMBER: DE 10043826.1  
/ PRIOR FILING DATE: 2000-09-01  
/ NUMBER OF SEQ ID NOS: 98  
/ SEQ ID NO 37  
/ LENGTH: 5501

; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)  
US-10-204-708-37

Query Match 9.3%; Score 39; DB 4; Length 5501;  
Best Local Similarity 49.8%; Pred. No. 0.26;  
Matches 126; Conservative 0; Mismatches 125; Indels 2; Gaps 1;  
Qy 102 GATTAGTTCTTCTTGGCCGATTAATCTTCTTCCGCGAAGATCTCCGTAAGAAG 161  
Db 1246 GTTGATGTAATATTTTAAATAATTTTGGGTTTTTGTGTGAATGTTATCGAGTTAATG 1305  
Qy 162 ATTTTAGTTAGAGATGAATCGTATAGTATGATTTTCAATCCAGATAGTTTCTTGTCTA 221  
Db 1306 TTATAGATAAGAGGTTTATTAATAATTTTGGGAATA-TTTATATTTTATTTTGA 1363  
Qy 222 GAATCTCGAAATCTCGATAGTTTTCACATGTGTAAATAGATTTCTTATTCGGCGAT 281  
Db 1364 TTTTATTTAGATGGTTGAATGAGTTTAAAGAGTTTTATGTTTTATTTTTCGTGTTATTA 1423  
Qy 282 TGTGATTAGGTTTTCATTTTCTTGTGATTATGCGATTGCAATTAGGCAATTTCTTTGGTT 341  
Db 1424 TATTAGTATATTTTATTTTATTTTGTGTTTAAAGAGATCGGGTTCGTATGTTGTTAGTT 1483  
Qy 342 TTGTTGATCTT 354  
Db 1484 TGGTTTGAATTT 1496

RESULT 10  
US-10-204-708-63  
; Sequence 63, Application US/10204708  
; Patent No. 6677731  
; GENERAL INFORMATION:  
; APPLICANT: OLEK, Alexander  
; APPLICANT: PIEPENBROCK, Christian  
; APPLICANT: BERLIN, Kurt  
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication  
; FILE REFERENCE: 5013.1012  
; CURRENT APPLICATION NUMBER: US/10/204,708  
; PRIOR FILING DATE: 2003-05-06  
; PRIOR APPLICATION NUMBER: PCT/EP01/03971  
; PRIOR FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: DE 10019058.8  
; PRIOR FILING DATE: 2000-04-06  
; PRIOR APPLICATION NUMBER: DE 10019173.8  
; PRIOR FILING DATE: 2000-04-07  
; PRIOR APPLICATION NUMBER: DE 10032529.7  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: DE 10043826.1  
; PRIOR FILING DATE: 2000-09-01  
; NUMBER OF SEQ ID NOS: 98  
; SEQ ID NO 63  
; LENGTH: 5562  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)  
US-10-204-708-63

Query Match 9.2%; Score 38.4; DB 4; Length 5562;  
Best Local Similarity 49.5%; Pred. No. 0.37;  
Matches 99; Conservative 0; Mismatches 101; Indels 0; Gaps 0;  
Qy 208 AGTTCTTGTCTAGAAATCTCTGAAATCTTCGATAGTTTTCACATGTGTAAATAGATGT 267  
Db 367 AGTTCTTGTCTAGAAATCTCTGAAATCTTCGATAGTTTTCACATGTGTAAATAGATGT 426  
Qy 268 TCTTATTCGGGATGTTGATAGGTTTTCATTTCTTGTGATTATGCGATTGCAATAGG 327

Db 427 TTTTATATAGTTTTTTTCGATTAGTGTTTTTTTTTTTGTTATTTAAAGTCGTTTTATTTT 486  
Qy 328 GATTTCCTTCTGTTTGTGTGATCTTACGATACATCTCTGCAATTCGATAGTATGAT 387  
Db 487 TTATTAGGTTTATTATTATTTTTTTTTTGTGATTGAAGAGTTTTTTTTTTAGTTTGTATAAAT 546  
Qy 388 CTAATCTTGTAAATTTGTT 407  
Db 547 ATTTTGTGTTTTTATTGTT 566

RESULT 11  
US-10-204-708-38  
; Sequence 38, Application US/10204708  
; Patent No. 6677731  
; GENERAL INFORMATION:  
; APPLICANT: OLEK, Alexander  
; APPLICANT: PIEPENBROCK, Christian  
; APPLICANT: BERLIN, Kurt  
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication  
; FILE REFERENCE: 5013.1012  
; CURRENT APPLICATION NUMBER: US/10/204,708  
; PRIOR FILING DATE: 2003-05-06  
; PRIOR APPLICATION NUMBER: PCT/EP01/03971  
; PRIOR FILING DATE: 2001-04-06  
; PRIOR APPLICATION NUMBER: DE 10019058.8  
; PRIOR FILING DATE: 2000-04-06  
; PRIOR APPLICATION NUMBER: DE 10019173.8  
; PRIOR FILING DATE: 2000-04-07  
; PRIOR APPLICATION NUMBER: DE 10032529.7  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: DE 10043826.1  
; PRIOR FILING DATE: 2000-09-01  
; NUMBER OF SEQ ID NOS: 98  
; SEQ ID NO 38  
; LENGTH: 5501  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)  
US-10-204-708-38

Query Match 9.1%; Score 38.2; DB 4; Length 5501;  
Best Local Similarity 49.7%; Pred. No. 0.42;  
Matches 97; Conservative 0; Mismatches 98; Indels 0; Gaps 0;  
Qy 210 TTTCTTTGTCTAGAAATCTCTGAAATCTTCGATAGTTTTCACATGTGTAAATAGATGTTTC 269  
Db 3630 TGTGTTTGTGTTAGTATGTTTTTTTTTATTGATGTTGTTTGTGTTGTTTGTGTTTGTGTTT 3689  
Qy 270 TTATTCGGCGATGTTGATTAGGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTTGTGTTT 329  
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Qy 390 AATCTTGTAAATTT 404  
Db 3810 ATATTATGTTATTTT 3824

RESULT 12  
US-10-204-708-61  
; Sequence 61, Application US/10204708  
; Patent No. 6677731  
; GENERAL INFORMATION:  
; APPLICANT: OLEK, Alexander  
; APPLICANT: PIEPENBROCK, Christian  
; APPLICANT: BERLIN, Kurt  
; TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication

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RESULT 13
US-10-204-708-20
  / Sequence 20, Application US/10204708
  / Patent No. 667731
  / GENERAL INFORMATION:
  / APPLICANT: OLEK, Alexander
  / APPLICANT: PIEPENEROCK, Christian
  / APPLICANT: BERLIN, Kurt
  / TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication
  / TITLE OF INVENTION: by Assessing DNA Methylation
  / FILE REFERENCE: 5013.1012
  / CURRENT APPLICATION NUMBER: US/10/204,708
  / CURRENT FILING DATE: 2003-05-06
  / PRIOR APPLICATION NUMBER: PCT/EP01/03971
  / PRIOR FILING DATE: 2001-04-06
  / PRIOR APPLICATION NUMBER: DE 10019058.8
  / PRIOR FILING DATE: 2000-04-06
  / PRIOR APPLICATION NUMBER: DE 10019173.8
  / PRIOR FILING DATE: 2000-04-07
  / PRIOR APPLICATION NUMBER: DE 10032529.7
  / PRIOR FILING DATE: 2000-06-30
  / PRIOR APPLICATION NUMBER: DE 10043826.1
  / PRIOR FILING DATE: 2000-09-01
  / NUMBER OF SEQ ID NOS: 98
  / SEQ ID NO 20
  / LENGTH: 6866
  / TYPE: DNA
  / ORGANISM: Artificial Sequence
  / FEATURE:
  / OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-204-708-20
Query Match          9.13; Score 38.2; DB 4; Length 6866;

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Query Match 9.1%; Score 38.2; DB 4; Length 6866;

/ TITLE OF INVENTION: Diagnosis of Diseases Associated with DNA Replication  
/ TITLE OF INVENTION: By Assessing DNA Methylation

/ FILE REFERENCE: 5013.1012  
/ CURRENT APPLICATION NUMBER: US/10/204,708  
/ CURRENT FILING DATE: 2003-05-06  
/ PRIOR APPLICATION NUMBER: PCT/EP01/03971  
/ PRIOR FILING DATE: 2001-04-06  
/ PRIOR APPLICATION NUMBER: DE 10019058.8  
/ PRIOR FILING DATE: 2000-04-06  
/ PRIOR APPLICATION NUMBER: DE 10019173.8  
/ PRIOR FILING DATE: 2000-04-07  
/ PRIOR APPLICATION NUMBER: DE 10032529.7  
/ PRIOR FILING DATE: 2000-06-30  
/ PRIOR APPLICATION NUMBER: DE 10043826.1  
/ PRIOR FILING DATE: 2000-09-01  
/ NUMBER OF SEQ ID NOS: 98  
/ SEQ ID NO 33  
/ LENGTH: 5455  
/ TYPE: DNA  
/ ORGANISM: Artificial Sequence  
/ FEATURE:  
/ OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)  
US-10-204-708-33

Query Match 9.0%; Score 37.8; DB 4; Length 5455;  
Best Local Similarity 53.8%; Pred. No. 0.54;  
Matches 78; Conservative 0; Mismatches 67; Indels 0; Gaps 0;  
  
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Db 3679 TTTTCTTTTGGGTTTTTTTATTTTTTTTGGGTTGGTAAATTAGT 3738  
  
QY 294 TTTTGATTTCCTGATTATGCGATTGCAATTAGGGATTTCCTTTGGTTTGTGTTGATCT 353  
Db 3739 TTTTAATTTTTTATATTTAGATTGGAAGTGGTGTTCATATTTTAGAGTT 3798  
  
QY 354 TAGATACATTCCTGCAATTGAATA 378  
Db 3799 GTTATTAATTTTTTTAAAGTTTTTA 3823

Search completed: June 20, 2004, 05:03:40  
Job time : 61.125 secs